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Recognition of double-stranded DNA using energetically activated duplexes with interstrand zippers of 1-, 2- or 4-pyrenyl-functionalized O2'-alkylated RNA monomers†

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Despite advances with triplex-forming oligonucleotides, peptide nucleic acids, polyamides and – more recently – engineered proteins, there remains an urgent need for synthetic ligands that enable specific recognition of double-stranded (ds) DNA to accelerate studies aiming at detecting, regulating and modifying genes. Invaders, *i.e.*, energetically activated DNA duplexes with interstrand zipper arrangements of intercalator-functionalized nucleotides, are emerging as an attractive approach toward this goal. Here, we characterize and compare Invaders based on 1-, 2- and 4-pyrenyl-functionalized O2'-alkylated uridine monomers **X–Z** by means of thermal denaturation experiments, optical spectroscopy, force-field simulations and recognition experiments using DNA hairpins as model targets. We demonstrate that Invaders with +1 interstrand zippers of **X** or **Y** monomers efficiently recognize mixed-sequence DNA hairpins with single nucleotide fidelity. Intercalator-mediated unwinding and activation of the double-stranded probe, coupled with extraordinary stabilization of probe–target duplexes (ΔT_m /modification up to +14.0 °C), provides the driving force for dsDNA recognition. In contrast, **Z**-modified Invaders show much lower dsDNA recognition efficiency. Thus, even very conservative changes in the chemical makeup of the intercalator-functionalized nucleotides used to activate Invader duplexes, affects dsDNA-recognition efficiency of the probes, which highlights the importance of systematic structure–property studies. The insight from this study will guide future design of Invaders for applications in molecular biology and nucleic acid diagnostics.

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Introduction

The right-handed DNA helix is one of the most fundamental structures in Nature due to its role as the carrier of genetic information. The two strands comprising the helix are held together through π – π stacking interactions between neighboring nucleobases and hydrogen bonding between base pairs.¹ Development of synthetic ligands that are capable of decoding the sequence information contained within double-stranded

DNA (dsDNA) has proven very challenging as the Watson–Crick base pairs are buried deeply within the duplex core and not readily accessible to exogenous agents. The most established dsDNA-targeting agents, *i.e.*, pyrrole–imidazole polyamides,² engineered proteins³ and triplex-forming oligonucleotides⁴ or peptide nucleic acids (PNAs),⁵ accordingly recognize chemical features that are accessible *via* one of the grooves instead.⁶ While these strategies are attractive, they are not without limitations. For example, PNA-based approaches normally require low salinity conditions, while triplex-based approaches require targets with polypurine tracts. Approaches with more relaxed sequence requirements have been developed but they still require the presence of purine stretches, a circumstance that may not be met at a target of interest.⁷ Other strategies allow recognition of mixed-sequence target regions that are unusually accessible to exogenous agents, such as AT-rich cruciforms or transcription bubbles.⁸

Synthetic ligands that recognize mixed-sequence B-DNA *via* duplex invasion are attractive due to the predictability of Watson–Crick base-pairing rules. However, these approaches

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†Electronic supplementary information (ESI) available: General experimental section; MS-data for modified ONs; representative thermal denaturation curves; additional denaturation and thermodynamic data; additional absorption and steady-state fluorescence emission spectra; complete data set of structural parameters from modeling studies; lowest energy representations of Type B structures; NMR spectra for nucleosides **2Y–4Z**. See DOI: 10.1039/c4ob01183j



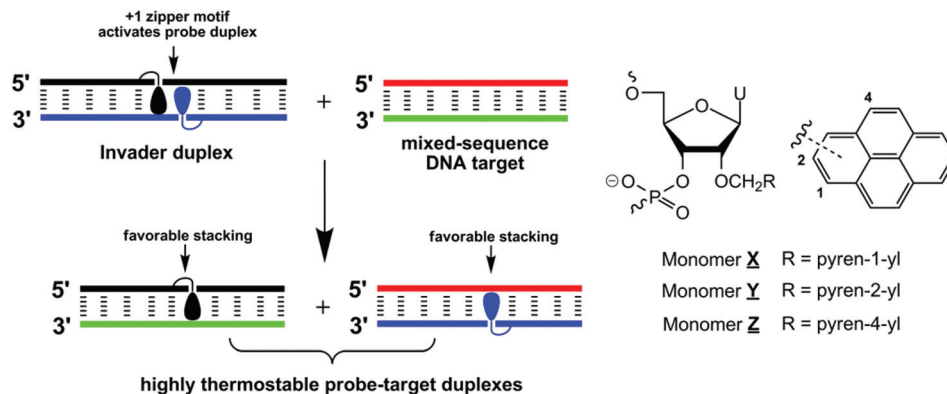


Fig. 1 Illustration of the Invader approach for recognition of mixed-sequence DNA and structures of monomers used herein. Droplets denote pyrene moieties.

must overcome a steep energetic penalty since pre-existing base pairs of *B*-DNA must be broken prior to probe binding. Pseudocomplementary DNA (pcDNA), *i.e.*, DNA duplexes featuring modified nucleobases that form weak base pairs with each other, while allowing hybridization to complementary DNA, recognize mixed-sequence target regions at *B*-DNA termini.⁹ This strategy has been successfully extended to pcPNA, which recognize internal regions of mixed-sequence dsDNA.¹⁰ However, the self-inhibitory effects observed at high pcPNA concentrations and the requirement for low salinity, pose potential limitations for pcPNA-mediated strand invasion in biological media.¹¹ γ -PNA, *i.e.*, single-stranded probes that are fully modified with conformationally pre-organized PNA building blocks, are another interesting class of dsDNA-targeting probes capable of recognizing mixed-sequence dsDNA targets. However, they too require non-physiological salinity for optimal strand invasion.¹² In summary, oligonucleotide-based probes that recognize mixed-sequence dsDNA targets at physiologic conditions remain largely elusive.

We have recently introduced *Invaders* as an alternative strategy toward mixed-sequence dsDNA recognition.¹³ These double-stranded probes are activated for dsDNA recognition through modification with +1 interstrand zippers of intercalator-functionalized nucleotides (for an illustration, see Fig. 1; for a formal definition of the zipper nomenclature, see the Experimental section). This structural motif results in a locally perturbed and destabilized region in the probe duplex since the motif represents a violation of the 'nearest neighbor exclusion principle', which states that intercalators, at most, bind to every second base pair of a DNA duplex due to limitations in local helix expandability.¹⁴ On the other hand, the two strands comprising an Invader probe display exceptionally strong affinity toward complementary single-stranded DNA (cDNA), since duplex formation results in strongly stabilizing pyrene-nucleobase stacking interactions (Fig. 1). In a key proof-of-concept study, we harnessed the energy difference between reactants (*i.e.*, Invader probes and target duplexes) and products (*i.e.*, probe-target duplexes) to drive recognition of chromosomal *DYZ-1* satellite

DNA in male bovine kidney cells at non-denaturing conditions.¹³

We originally used 2'-*N*-(pyren-1-yl)methyl-2'-amino- α -L-LNA (locked nucleic acid) nucleotides as the key activating components of Invader probes,¹⁵ but recently discovered that they can be replaced by 2'-*O*-(pyren-1-yl)methyl-ribonucleotides (Fig. 1).¹⁶ The resulting probes display similar dsDNA-recognition efficiency and are much easier to synthesize.^{17,18} Identification of these simple building blocks allowed us to initiate systematic structure-property relationship studies with the goal of gaining additional insights into the structural determinants governing Invader recognition efficiency. For example, we demonstrated that all four canonical 2'-*O*-(pyren-1-yl)methyl-RNA monomers can be used to construct dsDNA-binding Invader probes. However, probes using the pyrimidine monomers are particularly efficient.¹⁹

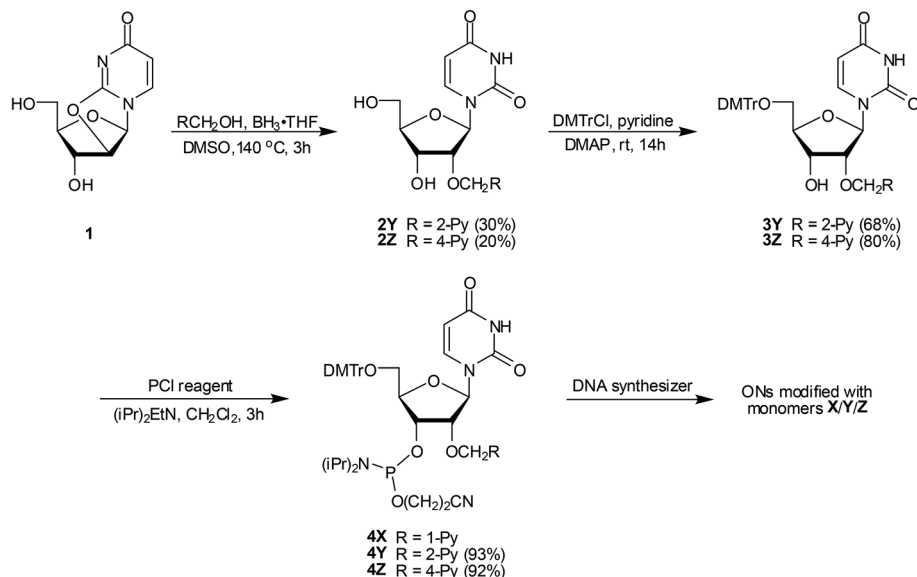
In the present study, we wanted to determine if the relative orientation between the pyrene intercalator and sugar skeleton has any impact on Invader-mediated recognition of mixed-sequence dsDNA. Toward this end, the corresponding 2- and 4-pyrenyl functionalized uridines were synthesized and incorporated into oligodeoxyribonucleotides (ONs), which were then characterized with respect to thermal denaturation, thermodynamic, UV-Vis absorption and fluorescence properties (Fig. 1). This was then followed by biophysical characterization of double-stranded probes with different interstrand arrangements of these monomers and recognition experiments using DNA hairpins as model targets.

Results and discussion

Synthesis of *O*2'-pyrene-functionalized uridine phosphoramidites

Phosphoramidites **4Y** and **4Z** were prepared in a similar manner as the 2'-*O*-(pyren-1-yl)methyl-uridine analogue **4X**.^{17b} Thus, treatment of *O*2,*O*2'-anhydrouridine **1**²⁰ with tris(pyren-2-yl)methyl borate or tris(pyren-4-yl)methyl borate – generated *in situ*²¹ via addition of 2-pyrenemethanol²² or 4-pyrenemetha-





Scheme 1 Synthesis of O2'-pyrene-functionalized uridine phosphoramidites **4Y** and **4Z**. DMTr = 4,4'-dimethoxytrityl; PCI reagent = 2-cyanoethyl-*N,N'*-diisopropylchlorophosphoramidite; 1-Py = pyren-1-yl; 2-Py = pyren-2-yl; 4-Py = pyren-4-yl.

mol²³ to borane – afforded **2Y** and **2Z** in modest but acceptable yields (30% and 20% respectively, Scheme 1). Subsequent O5'-dimethoxytritylation gave nucleosides **3Y** and **3Z**, which upon treatment with 2-cyanoethyl-*N,N'*-diisopropylchlorophosphoramidite (PCI reagent) and Hünig's base provided target phosphoramidites **4Y** and **4Z** in excellent yield.

Synthesis of modified ONs and experimental design

Phosphoramidites **4X**, **4Y** and **4Z** were used in machine-assisted solid-phase DNA synthesis to incorporate monomers **X–Z** into ONs using 4,5-dicyanoimidazole as an activator and extended hand-coupling (15 min), which resulted in stepwise coupling yields of ~99%/~99%/~98% for **4X/4Y/4Z**, respectively. The identity and purity of the modified ONs was established *via* MALDI-TOF (Table S1 in the ESI†) and ion-pair reverse phase HPLC (>85% purity), respectively.

Monomers **X–Z** were studied in 9-mer sequence contexts, which we have previously used to screen and identify potential Invader building blocks.¹⁶ ONs containing a single incorporation in the 5'-GBG ATA TGC context are denoted **X1**, **Y1**, and **Z1**. Similar conventions apply for the **B2–B6** series (Table 1). Reference DNA and RNA strands are denoted **D1/D4** and **R1/R4**, respectively (see footnote, Table 1).

Thermostability of duplexes with complementary DNA/RNA

First, the thermostabilities of duplexes between **B1–B6** and complementary DNA or RNA targets were determined from thermal denaturation experiments performed in medium salt phosphate buffer ([Na⁺] = 110 mM, pH 7.0). The resulting denaturation curves display the expected monophasic sigmoidal transitions (Fig. S1 in the ESI†). Interestingly, **Y**-modified ONs form much more thermostable duplexes with cDNA than

Table 1 Thermal denaturation temperatures (*T_m*'s) for duplexes between **B1–B6** and complementary DNA or RNA^a

ON	Sequence	B =	<i>T_m</i> (Δ <i>T_m</i> /mod) [°C]					
			+cDNA			+cRNA		
			X ^b	Y	Z	X ^b	Y	Z
B1	5'-GBG ATA TGC		34.5 [+5.0]	35.5 [+6.0]	29.5 [±0.0]	24.5 [−2.0]	25.5 [−1.5]	20.5 [−6.0]
B2	5'-GTG ABA TGC		42.5 [+13.0]	43.5 [+14.0]	37.5 [+8.0]	30.5 [+4.0]	34.0 [+7.5]	27.5 [+1.0]
B3	5'-GTG ATA BGC		37.5 [+8.0]	39.0 [+9.5]	33.5 [+4.0]	26.5 [±0.0]	28.0 [+1.5]	21.5 [−5.0]
B4	3'-CAC BAT ACG		33.0 [+3.5]	35.5 [+6.0]	26.5 [−3.0]	20.5 [−4.5]	23.0 [−1.5]	16.5 [−8.0]
B5	3'-CAC TAB ACG		42.5 [+13.0]	43.5 [+14.0]	38.5 [+9.0]	27.5 [+2.5]	32.0 [+7.5]	26.5 [+2.0]
B6	3'-CAC BAB ACG		43.5 [+7.0]	45.5 [+8.0]	32.5 [+1.5]	24.0 [−0.3]	28.5 [+2.0]	16.5 [−4.0]

^a Δ*T_m* = change in *T_m* relative to reference duplexes **D1:D4** (*T_m* ≡ 29.5 °C), **D1:R4** (*T_m* ≡ 26.5 °C) or **R1:D4** (*T_m* ≡ 24.5 °C), where **D1**: 5'-GTGATATGC, **D4**: 3'-CACTATACG, **R1**: 5'-GUGAUAUGC and **R4**: 3'-CACUAUACG; *T_m*'s are determined as the maximum of the first derivative of melting curves (*A*₂₆₀ vs. *T*) recorded in medium salt buffer ([Na⁺] = 110 mM, [Cl[−]] = 100 mM, pH 7.0 (NaH₂PO₄/Na₂HPO₄)), using 1.0 μM of each strand. *T_m*'s are averages of at least two measurements within 1.0 °C; A = adenin-9-yl DNA monomer, C = cytosin-1-yl DNA monomer, G = guanin-9-yl DNA monomer, T = thymine-1-yl DNA monomer. For structures of monomers **X–Z** see Fig. 1. ^b From ref. 17b.



Table 2 Discrimination of mismatched DNA targets by **X2/Y2/Z2** and reference strands^a

ON	Sequence	B =	DNA: 3'-CAC TBT ACG			
			T_m [°C]		ΔT_m [°C]	
			A	C	G	T
D1	5'-GTG ATA TGC		29.5	-16.5	-9.5	-17.0
X2 ^b	5'-GTG AXA TGC		42.5	-13.5	-5.5	-7.0
Y2	5'-GTG AYA TGC		43.5	-16.5	-6.5	-9.0
Z2	5'-GTG AZA TGC		37.5	-10.0	-4.0	-3.0

^a For conditions of thermal denaturation experiments, see Table 1. T_m 's of fully matched duplexes are shown in bold. ΔT_m = change in T_m relative to fully matched duplex. ^b From ref. 17b.

unmodified ONs ($\Delta T_m/\text{mod} = +6.0$ to $+14.0$ °C, Table 1). In fact, the resulting duplexes are even more thermostable than **X**-modified duplexes, suggesting that the 2-pyrenyl moiety of monomer **Y** is very well accommodated in DNA duplexes. In contrast, **Z**-modified ONs display considerably lower affinity toward cDNA ($\Delta T_m/\text{mod} = -3.0$ to $+9.0$ °C, Table 1). The thermostability trends are sequence-dependent. Thus, ONs in which the pyrene-functionalized monomers are flanked by 3'-purines, induce greater stabilization than ONs with 3'-flanking pyrimidines (e.g., compare $\Delta T_m/\text{mod}$ values for **B2**- and **B4**-series, Table 1). This is indicative of 3'-intercalative binding modes of the pyrene moieties,^{19,24} as 3'-flanking purines would provide larger π -stacking surfaces than 3'-pyrimidines. Less stable duplexes are formed with cRNA ($\Delta T_m/\text{mod} = -8.0$ to $+7.5$ °C, Table 1; trend: **Y** > **X** > **Z**), which also points to intercalative pyrene binding modes,^{18,24a,25} as intercalators generally favor the less compressed B-type helix geometry of DNA: DNA duplexes.²⁶

Binding specificity

Next, the binding specificity of singly modified ONs (**B2**-series) was studied using DNA targets with mismatched nucleotides opposite of the pyrene-functionalized monomer (Table 2). **X2/Y2/Z2** discriminate mismatched DNA targets less efficiently than reference strand **D1**, especially when a mismatched T is opposite of the modification. This is in agreement with other

studies in which reduced binding specificity of intercalator-modified ONs was observed.²⁶ Interestingly, the specificity trends mirror the affinity trends. Thus, **Y2** discriminates mismatched target more efficiently than **X2**, which, conversely, discriminates mismatched targets more efficiently than **Z2**. Additional specificity data are discussed in the ESI (Tables S3 and S4†).

Optical spectroscopy

UV-Vis absorption and steady-state fluorescence emission spectra of **X/Y/Z**-modified ONs in the presence or absence of complementary DNA/RNA targets were recorded next to further ascertain intercalative binding modes of the pyrene moieties, as intercalation is known to induce bathochromic shifts of pyrene absorption peaks²⁷ and nucleobase-mediated quenching of pyrene fluorescence.^{27a,28} Indeed, **X/Y/Z**-modified ONs generally display hypochromic and bathochromic shifts in the pyrene absorption spectra upon hybridization with DNA and RNA targets ($\Delta\lambda_{\text{max}} = 0$ –5 nm, Table 3; Fig. S2–S4 in the ESI†). Slightly greater bathochromic shifts are generally observed upon hybridization with DNA than RNA targets, which again may reflect the preference of intercalators for the less compressed geometry of B-DNA.

Steady-state fluorescence emission spectra of **X/Y/Z**-modified ONs and the corresponding duplexes with cDNA/cRNA display two vibronic bands at $\lambda_{\text{em}} = 382 \pm 3$ nm and 402 ± 3 nm, respectively, as well as a small shoulder at ~ 420 nm. As anticipated, the fluorescence intensity typically decreases upon hybridization with DNA/RNA targets, with greater decreases typically being observed upon DNA binding (Fig. 2; Fig. S5–S7 in the ESI†). Interestingly, duplexes modified with monomer **Z** are noticeably less fluorescent than **X**- or **Y**-modified duplexes.

Molecular modeling of probe–target duplexes

To rationalize the observed biophysical trends, we performed force-field calculations on duplexes between **X2/Y2/Z2** and complementary DNA (residue numbering: 5'-G₁T₂G₃A₄B₅A₆-T₇G₈C₉-3'-C₁₈A₁₇C₁₆T₁₅A₁₄T₁₃A₁₂C₁₁G₁₀). The complete data set of structural furanose, base pair and dinucleotide step para-

Table 3 Absorption maxima in the 335–355 nm region for single-stranded **X/Y/Z**-modified ONs and the corresponding duplexes with complementary DNA or RNA^a

ON	Sequence	B =	λ_{max} [$\Delta\lambda_{\text{max}}$]/nm								
			X ^b			Y			Z		
			SSP	+cDNA	+cRNA	SSP	+cDNA	+cRNA	SSP	+cDNA	+cRNA
B1	5'-GBG ATA TGC		350	353[+3]	352[+2]	344	346[+2]	345[+1]	343	344[+1]	343[±0]
B2	5'-GTG ABA TGC		348	353[+5]	352[+4]	342	345[+3]	345[+3]	340	343[+3]	343[+3]
B3	5'-GTG ATA BGC		350	353[+3]	352[+2]	344	346[+2]	345[+1]	343	344[+1]	343[±0]
B4	3'-CAC BAT ACG		350	352[+2]	352[+2]	344	345[+1]	345[+1]	342	344[+2]	343[+1]
B5	3'-CAC TAB ACG		349	353[+4]	352[+3]	343	345[+2]	345[+2]	342	344[+2]	344[+2]

^a Measurements were performed at 5 °C using a spectrophotometer and quartz optical cells with 1.0 cm path lengths. Buffer conditions are as for thermal denaturation experiments. ^b From ref. 17b.



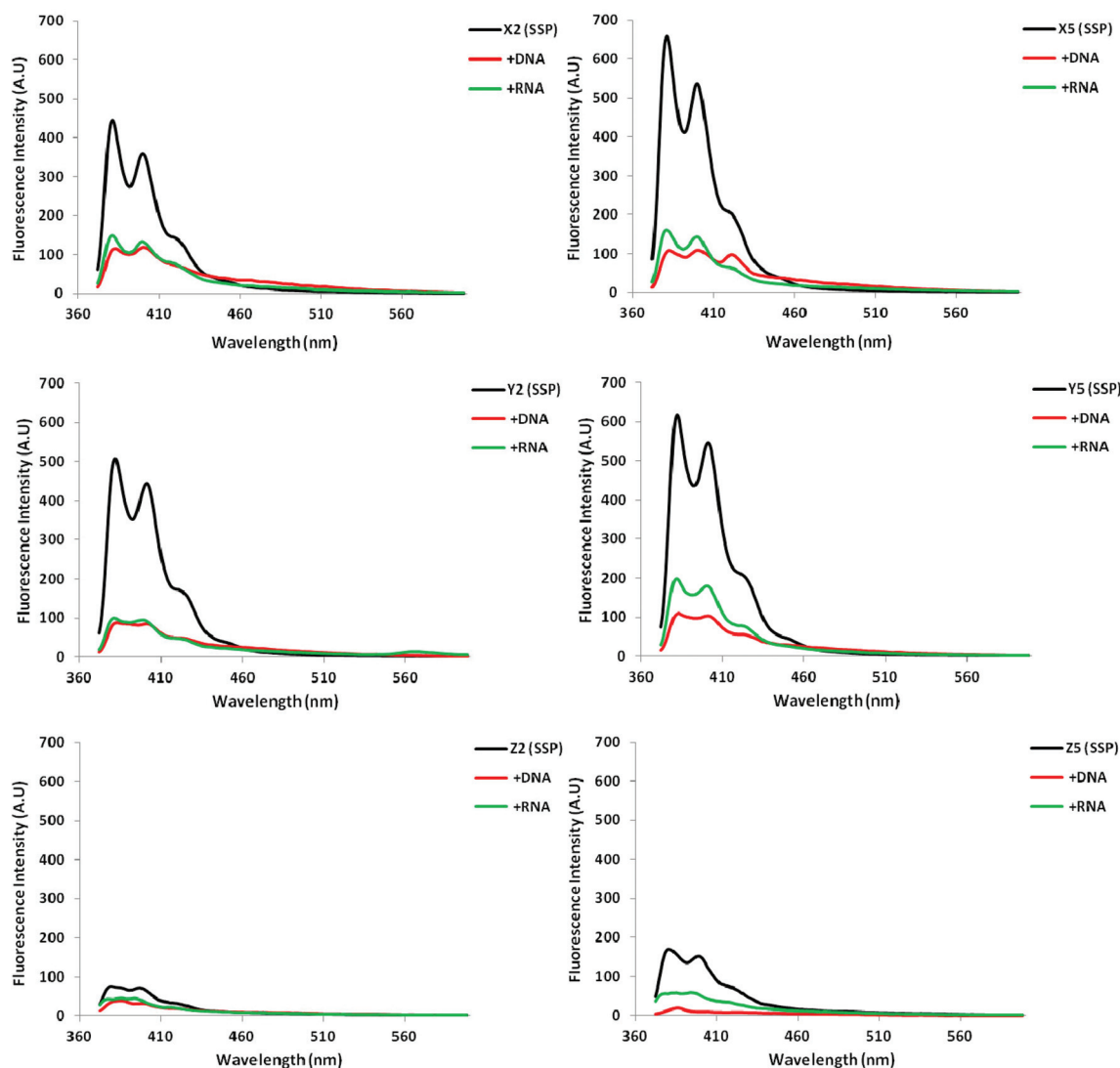


Fig. 2 Steady-state fluorescence emission spectra of select X/Y/Z-modified ONs and corresponding duplexes with DNA/RNA targets. Spectra were recorded at $T = 5^\circ\text{C}$ using $\lambda_{\text{ex}} = 350, 345$ and 340 nm for X, Y and Z-modified ONs, respectively. Each strand was used at $1.0\text{ }\mu\text{M}$ concentration in T_m buffer.

meters²⁹ is provided in the ESI (Tables S7–S8 and Fig. S9–S14†).

In agreement with the photophysical data presented above, the pyrene moieties remain stably intercalated in the duplex core throughout the stochastic dynamics simulations (Fig. 3 top – for details of calculation protocol, see Experimental section). Inspection of the structures reveals that the right-handed duplexes are elongated relative to the corresponding B-DNA reference duplex (e.g., rise increases from $\sim 3.3\text{ }\text{\AA}$ to $\sim 7.3\text{ }\text{\AA}$ across the $B_5A_6:A_{14}T_{13}$ dinucleotide step, Fig. S10 in the ESI†). The overlap between the pyrene and neighboring nucleobases is strongly influenced by the substitution pattern of the pyrene, which explains the observed thermostability trends (T_m decreases $Y \geq X > Z$). The pyrene of monomer X stacks with the nucleobase moieties of X_5 , A_6 and A_{13} , while the pyrene of monomer Y is positioned in a manner that facili-

tates stacking with all four neighboring nucleobase moieties (Fig. 3 bottom). Conversely, the pyrene of monomer Z stacks only with the flanking nucleobases on its own strand, i.e., Z_5 and A_6 . The poorer fit of the pyrene of monomer Z is also reflected in greater perturbation across the $B_5A_6:A_{14}T_{13}$ dinucleotide step, i.e., more pronounced *buckle* ($\sim 6^\circ$ and $\sim 4^\circ$ in **D1:D4**, $\sim 13^\circ$ and $\sim 10^\circ$ in **X2:D4**, $\sim 15^\circ$ and $\sim 11^\circ$ in **Y2:D4**, and $\sim 21^\circ$ and $\sim 19^\circ$ in **Z2:D4**, Fig. S9 in the ESI†) and decreased *roll* ($\sim 12^\circ$ in **D1:D4**, $\sim 10^\circ$ in **X2:D4**, $\sim 11^\circ$ in **Y2:D4**, and $\sim 19^\circ$ in **Z2:D4**, Fig. S10 in the ESI†). Another noteworthy observation is that the furanose rings of nucleotides X_5A_6 , Y_5A_6 and Z_5A_6 generally exhibit increased *South* type character relative to the reference duplex ($P = 124\text{--}171^\circ$ vs. $102\text{--}129^\circ$), presumably as a consequence of π - π -stacking between the pyrene and flanking nucleobases (Table S7 in the ESI†).



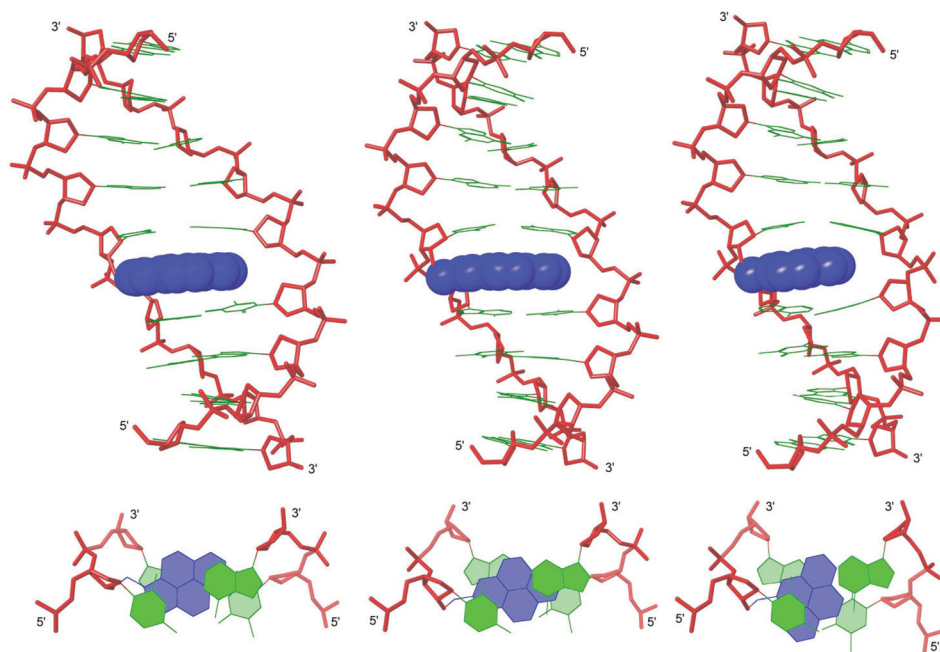


Fig. 3 Lowest energy structures of X2:D4 (left), Y2:D4 (middle) and Z2:D4 (right). Upper: side view of duplex; lower: top view of the central duplex region. Color code: sugar phosphate backbone (red); pyrene moieties (blue) and nucleobases (green). Hydrogen atoms, sodium ions and bond orders are omitted for clarity.

To sum up, the molecular modeling structures are useful in explaining the observed thermostability trends and photo-physical properties of X/Y/Z-modified duplexes.

Biophysical properties of duplexes with interstrand zippers of X/Y/Z-monomers

Next, we determined the thermostability of DNA duplexes with different interstrand zipper arrangements of X/Y/Z monomers to identify monomers and probe architectures that show the greatest potential for dsDNA-recognition. The impact on duplex thermostability upon incorporation of a second monomer can be additive, greater-than-additive or less-than-additive relative to the corresponding singly modified duplex. The impact is conveniently approximated in terms of T_m 's by the term 'deviation from additivity' (DA) defined as: $DA_{\text{ONA:ONB}} \equiv \Delta T_m (\text{ON}_A:\text{ON}_B) - [\Delta T_m (\text{ON}_A:\text{cDNA}) + \Delta T_m (\text{cDNA}:\text{ON}_B)]$, where $\text{ON}_A:\text{ON}_B$ is a duplex with an interstrand arrangement of monomers. DA also serves as an indicator for dsDNA-recognition potential. Probes with strongly negative DA values are likely to be activated for recognition of iso-sequential dsDNA *via* the process depicted in Fig. 1, since the products of the recognition process (*i.e.*, probe-target duplexes) are more thermostable than the reactants (*i.e.*, probe duplexes and target duplexes). A more rigorous approach based on differences in ΔG values of probe-target and Invader duplexes is discussed shortly.

As expected,^{13,15,16} duplexes with +1 interstrand monomer arrangements are less thermostable and more strongly activated for dsDNA-recognition than duplexes with other arrangements (compare T_m 's and DA values for B2:B5 relative to other

probe duplexes, Table 4). Clearly the two pyrene-functionalized monomers interact with each other in an energetically unfavorable manner when placed in this motif. Interestingly, there is little difference in the dsDNA-targeting potential of X2:X5, Y2:Y5 and Z2:Z5 as judged by the DA values; the lower thermostability of Z-modified probe-target duplexes is compensated by an equivalent destabilization of the probe duplexes (compare DA values for X2:X5, Y2:Y5 and Z2:Z5, Table 4).

The above T_m -based conclusions were corroborated through analysis of thermodynamic parameters for duplex formation, which were derived through fitting of thermal denaturation curves.³⁰ Thus, formation of probe-target duplexes is highly favorable relative to unmodified duplexes, with Y-modified and Z-modified ONs leading to the most and least stable duplexes with cDNA, respectively (see the first two $\Delta\Delta G^{293}$ columns, Table 4). Stabilization of the probe-target duplexes is largely a consequence of increased entropy (Tables S5 and S6 in the ESI†). On the other hand, B2:B5 duplexes are much less stable (see the third $\Delta\Delta G^{293}$ column, Table 4) due to strongly increased enthalpy (Tables S5 and S6 in the ESI†). Consequentially, B2:B5 probes display favorable binding free energy for recognition of iso-sequential dsDNA targets as given by $\Delta G_{\text{rec}}^{293} (\text{ON}_A:\text{ON}_B) = \Delta G^{293} (\text{ON}_A:\text{cDNA}) + \Delta G^{293} (\text{cDNA}:\text{ON}_B) - \Delta G^{293} (\text{ON}_A:\text{ON}_B) - \Delta G^{293} (\text{dsDNA})$ (*i.e.*, $\Delta G_{\text{rec}}^{293}$ for B2:B5 $\ll 0$ kJ mol⁻¹, Table 4). X2:X5 and Y2:Y5 display very similar $\Delta G_{\text{rec}}^{293}$ values. Unfortunately, the absence of a clear lower base line in the thermal denaturation curve of Z2:Z5 precluded determination of $\Delta G_{\text{rec}}^{293}$ for this duplex. Probes with other interstrand arrangements of X/Y/Z-monomers are much more stable and display much less favorable binding free energy for recognition



Table 4 Properties of X/Y/Z-modified probe duplexes^a

ON	ZP	Sequence	T_m (°C)	DA (°C)	ΔG^{293} [$\Delta\Delta G^{293}$] (kJ mol ⁻¹)		Probe duplex	ΔG_{rec}^{293} (kJ mol ⁻¹)	λ_{max} (nm)
					Upper ON vs. cDNA	Lower ON vs. cDNA			
X1	+4	5'-G X G ATA TGC	47.5	± 0.0	-46 ± 0 [-5]	-52 ± 0 [-11]	-55 ± 1 [-14]	-2	352
X5		3'-CAC T A X ACG							
X1	+2	5'-G X G ATA TGC	31.5	-6.5	-46 ± 0 [-5]	-46 ± 0 [-5]	-45 ± 1 [-4]	-6	350
X4		3'-CAC X AT ACG							
X2	+1	5'-GTG A X A TGC	26.5	-29.0	-55 ± 1 [-14]	-53 ± 1 [-12]	-38 ± 1 [+3]	-29	348
X5		3'-CAC T A X ACG							
X2	-1	5'-GTG A X A TGC	39.5	-6.5	-54 ± 0 [-13]	-46 ± 0 [-5]	-51 ± 1 [-10]	-8	351
X4		3'-CAC X AT ACG							
X3	-1	5'-GTG ATA X GC	45.5	-5.0	-50 ± 0 [-9]	-52 ± 0 [-11]	-55 ± 1 [-14]	-6	352
X5		3'-CAC T A X ACG							
X3	-3	5'-GTG ATA X GC	40.5	-0.5	-50 ± 0 [-7]	-46 ± 0 [-3]	-53 ± 1 [-10]	± 0	352
X4		3'-CAC X AT ACG							
Y1	+4	5'-G Y G ATA TGC	48.5	-1.0	-48 ± 0 [-7]	-55 ± 1 [-14]	-56 ± 1 [-15]	-6	346
Y5		3'-CAC T A Y ACG							
Y1	+2	5'-G Y G ATA TGC	35.5	-6.0	-48 ± 0 [-7]	-46 ± 0 [-5]	-45 ± 1 [-4]	-8	346
Y4		3'-CAC Y AT ACG							
Y2	+1	5'-GTG A Y A TGC	28.5	-29.5	-56 ± 0 [-15]	-55 ± 1 [-14]	-43 ± 1 [-2]	-27	339
Y5		3'-CAC T A Y ACG							
Y2	-1	5'-GTG A Y A TGC	43.5	-6.0	-56 ± 0 [-15]	-46 ± 0 [-5]	-55 ± 1 [-14]	-6	346
Y4		3'-CAC Y AT ACG							
Y3	-1	5'-GTG ATA Y GC	47.5	-5.5	-51 ± 1 [-10]	-55 ± 1 [-14]	-58 ± 1 [-17]	-7	346
Y5		3'-CAC T A Y ACG							
Y3	-3	5'-GTG ATA Y GC	43.0	-2.0	-51 ± 1 [-10]	-46 ± 0 [-5]	-53 ± 1 [-12]	-3	346
Y4		3'-CAC Y AT ACG							
Z1	+4	5'-G Z G ATA TGC	38.5	± 0.0	-44 ± 0 [-3]	-48 ± 1 [-7]	-49 ± 1 [-8]	-2	345
Z5		3'-CAC T A Z ACG							
Z1	+2	5'-G Z G ATA TGC	27.5	+1.0	-44 ± 0 [-3]	-41 ± 0 [± 0]	-41 ± 1 [± 0]	-3	342
Z4		3'-CAC Z AT ACG							
Z2	+1	5'-GTG A Z A TGC	19.5	-27.0	-48 ± 1 [-7]	-48 ± 1 [-7]	N/A	N/A	339
Z5		3'-CAC T A Z ACG							
Z2	-1	5'-GTG A Z A TGC	30.5	-4.0	-48 ± 1 [-7]	-41 ± 0 [± 0]	-41 ± 1 [± 0]	-7	344
Z4		3'-CAC Z AT ACG							
Z3	-1	5'-GTG ATA Z GC	37.0	-5.5	-46 ± 0 [-5]	-48 ± 1 [-7]	-46 ± 1 [-5]	-7	345
Z5		3'-CAC T A Z ACG							
Z3	-3	5'-GTG ATA Z GC	32.5	+2.0	-46 ± 0 [-5]	-41 ± 0 [± 0]	-42 ± 1 [-1]	-4	345
Z4		3'-CAC Z AT ACG							

^a ZP = zipper. For conditions of thermal denaturation and absorption experiments, see Tables 1 and 3, respectively. DA = ΔT_m (Invader) - (ΔT_m (upper strand vs. cDNA) + ΔT_m (lower strand vs. cDNA)). $\Delta\Delta G^{293}$ is measured relative to ΔG^{293} for **D1:D4** = -41 kJ mol⁻¹. ΔG_{rec}^{293} = ΔG^{293} (upper strand vs. cDNA) + ΔG^{293} (lower strand vs. cDNA) - ΔG^{293} (probe duplex) - ΔG^{293} (dsDNA target). " \pm " denotes standard deviation. N/A = the absence of a clear lower base line prevented determination of this value. T_m 's for X-modified duplexes have been previously published in ref. 17b but are included to facilitate direct comparison.

of iso-sequential dsDNA targets (ΔG_{rec}^{293} between -8 and 0 kJ mol⁻¹, Table 4), presumably since the two monomers act independently from each other.

The unique characteristics of duplexes with +1 interstrand monomer arrangements relative to other probe duplexes are also reflected in the blue-shifted pyrene absorption peaks (Fig. S8 in the ESI†), which are indicative of reduced pyrene-nucleobase interactions (compare λ_{max} for **B2:B5** with λ_{max} for other probe duplexes (Table 4) or probe-target duplexes (Table 3)).

All probe duplexes exhibit vibronic peaks at λ_{em} = 380 \pm 2 nm, 400 \pm 3 nm and \sim 420 \pm 3 nm in their steady-state fluorescence emission spectrum (Fig. 4). In addition, duplexes with +2 interstrand monomer arrangements display prominent

and unstructured emission centered at λ_{em} \sim 490 nm, which is consistent with pyrene excimer emission (Fig. 4).^{15,16,19,31} The presence of these signals shows that the pyrene moieties are stacking with an interplanar separation of \sim 3.4 Å. We speculate that stacking occurs in the major groove in a similar manner as previously suggested for duplexes with +2 zipper arrangements of pyrene-functionalized *ara*-uridine monomers.^{31b,32} Duplexes with +1 interstrand monomer arrangements also display excimer emission albeit of much weaker intensity. In this case, the two pyrene moieties most likely stack inside the duplex core in a similar manner as recently suggested for duplexes modified with 2'-N-(pyren-1-yl)methyl-2'-amino- α -L-LNA monomers; the spatial separation for



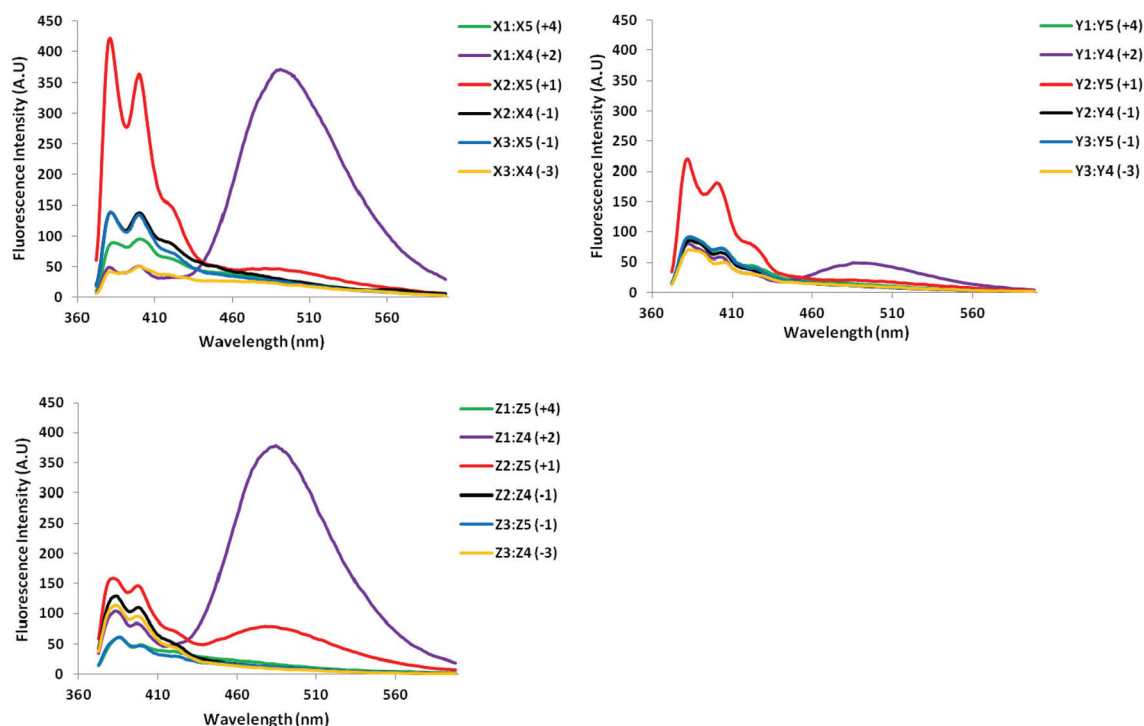


Fig. 4 Steady-state fluorescence emission spectra of duplexes with different interstrand monomer arrangements of X, Y, and Z (zipper type indicated in parenthesis). For experimental conditions, see Fig. 2. Spectra for X-modified duplexes were previously reported in ref. 16 but are included for comparison.

pyrene–pyrene stacking in the major groove is too large.^{15a,16} Dual intercalation would also explain the energetic lability of the **B2:B5** duplexes, as a localized region with one intercalator per base pair would ensue, which represents a violation of the ‘nearest neighbor exclusion principle’.¹⁴

Molecular modeling of B2:B5 duplexes

Force-field based simulations of **B2:B5** duplexes were performed to gain additional insight into the binding mode of the pyrene moieties. The results of these simulations point toward two possible binding modes, *i.e.*, Type A in which both pyrene moieties intercalate into the duplex core (Fig. 5) and Type B in which one pyrene is intercalating and the other is extruded into the major groove (Fig. S15 in the ESI†). The presence of excimer signals in the steady state emission spectra of **X2:X5** and **Z2:Z5** (Fig. 4), suggests that these duplexes are capable of adopting Type A conformations. However, the weak intensity of the excimer signals – especially with **Y2:Y5** – suggests that **B2:B5** duplexes equilibrate between Type A and Type B conformations. Type A structures are characterized by significant elongation (*rise* ~ 10 Å) and local perturbation (*buckle* $\sim 30^\circ$ and $\sim -30^\circ$ at B_5A_6 and $A_{14}B_{13}$, respectively, and *roll* $\sim -30^\circ$ across X_5A_6 : $X_{14}B_{13}$ and Y_5A_6 : $Y_{14}B_{13}$ and $\sim -18^\circ$ across Z_5A_6 : A_{14}) (Fig. S11–S12 in the ESI†). Type B structures are less elongated (*rise* ~ 7.5 Å) and less perturbed (*buckle* $\sim 20^\circ$ and $\sim -20^\circ$ at B_5A_6 and $A_{14}B_{13}$, respectively, and *roll* $\sim -5^\circ$) (Fig. S13–S14 in the ESI†). Both conformations can account for the lability of the **B2:B5**

duplexes; base pairing in the vicinity of the interstrand zipper is distorted in both structures and the presence of a large hydrophobic pyrene in the major groove (Type B) will almost certainly perturb the stabilizing hydration layer or cause unfavorable steric interactions.

Recognition of DNA hairpins using activated probe duplexes

The $\Delta G_{\text{rec}}^{293}$ values indicate that duplexes with +1 interstrand zippers of **X/Y/Z** monomers are the most strongly activated probes for dsDNA-recognition (Table 4). We therefore decided to evaluate these probes in greater detail using a gel shift assay that we have used to screen other potential Invader monomers with.^{13,16} Thus, a digoxigenin (DIG) labeled DNA hairpin (DH) – comprised of a 9-mer double-stranded mixed-sequence stem that is linked by a T_{10} loop – was used as a model target (Fig. 6a).³³ As expected, the unimolecular nature of **DH1** strongly stabilizes the stem region as seen from the significantly higher T_m relative to the corresponding linear DNA duplex ($T_m = 57.0$ °C vs. 29.5 °C, respectively; Fig. 6b and Table 1). Incubation of **DH1** with **X2:X5** or **Y2:Y5** in a HEPES buffer for 15 h at ambient temperature resulted in dose-dependent recognition as evidenced by the emergence of a slower migrating band on non-denaturing PAGE gels (Fig. 6c).³³ It is particularly noteworthy that as little as 1.0 molar equivalent of **Y2:Y5** resulted in $\sim 50\%$ dsDNA-recognition (Fig. 6d). In contrast, very little recognition was observed even at 500-fold molar excess of **Z2:Z5** (Fig. 6c and 6d). This was surprising to us considering the very similar DA values of the **B2:B5**



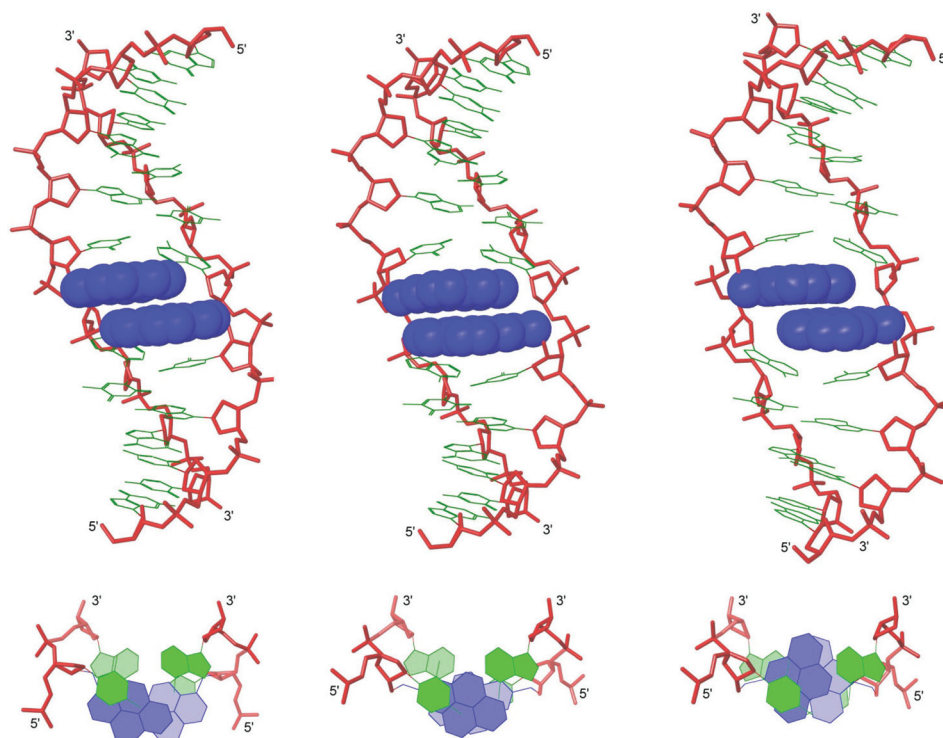


Fig. 5 Lowest energy structures of X2:X5 (left), Y2:Y5 (middle) and Z2:Z5 (right) in Type A conformation. Top: side view of duplex; bottom: top view of the central duplex region. For color code, see Fig. 3.

duplexes (Table 4). To determine if the poor dsDNA-recognition efficiency of Z2:Z5 was due to insufficient stability of the recognition complex and/or slow reaction kinetics, we performed separate experiments in which DH1 was annealed in the presence of 500-fold molar excess of X2:X5 or Z2:Z5 (*i.e.*, heated to 95 °C for 2 min, cooled to 8° over 3 h, then analyzed on non-denaturing PAGE gels). The resulting gel electrophoretograms (results not shown) were essentially identical to those from the room temperature incubation experiments discussed above, which suggests that the recognition complex between Z2:Z5 and DH1 is not sufficiently stable at the experimental conditions of the assay.

As expected, the unmodified control duplex D1:D4 did not result in formation of recognition complexes even when used at 500-fold molar excess, as there is no driving force to overcome the energetic penalty of opening DH1 (Fig. 6c). The use of 500-fold molar excess of single-stranded X2/X5/Y2/Y5 only resulted in 15–40% recognition of DH1 (Fig. 6e), which demonstrates that both probe strands are necessary for efficient dsDNA-recognition.

Finally, the binding specificity of X2:X5 and Y2:Y5 was examined in detail by incubating the probe duplexes with DNA hairpins DH2–DH7, which are fully base-paired but singly mismatched relative to the Invader probes (underlined residues indicate position of sequence deviation, Fig. 6b). Importantly, neither a 500-fold molar excess of X2:X5 nor Y2:Y5 resulted in recognition of the singly mismatched DNA hairpins, which demonstrates that hairpin recognition is highly specific (Fig. 6f).

Conclusion

Short synthetic routes to suitably protected 2'-O-(pyren-2-yl)-methyluridine and 2'-O-(pyren-4-yl)methyluridine have been developed. ONs modified with 2'-O-(pyren-2-yl)methyluridines display greater affinity toward complementary DNA and better mismatch discrimination than ONs modified with the corresponding 1-pyrenyl or 4-pyrenyl analogues. Molecular modeling suggests this to be a consequence of more efficient π - π -stacking between the intercalating pyrene moiety and neighboring base pairs.

DNA duplexes with +1 interstrand arrangements of these pyrene-functionalized monomers display lower thermostability, more blue-shifted pyrene absorption maxima, more distinctive fluorescence emission profiles and greater potential for dsDNA-recognition than duplexes with other monomer arrangements. Force field simulations suggest that the +1 zipper motif results in significant perturbation of the duplex. Although the chemical differences between the three studied monomers are relatively minor, significant variations in the dsDNA-recognition efficiencies of the resulting probes are observed. Thus, probes with +1 interstrand zippers of 2'-O-(pyren-2-yl)methyluridines recognize mixed-sequence DNA hairpins very efficiently, whereas probes constructed with the corresponding 2'-O-(pyren-4-yl)methyluridines do not. Remarkably, Invaders based on 2'-O-(pyren-2-yl)methyluridines result in ~50% recognition of dsDNA when used in equimolar quantities. These findings encourage additional structure–property relationship studies based on the 2-pyrenyl scaffolds with the goal of further increasing the dsDNA recog-



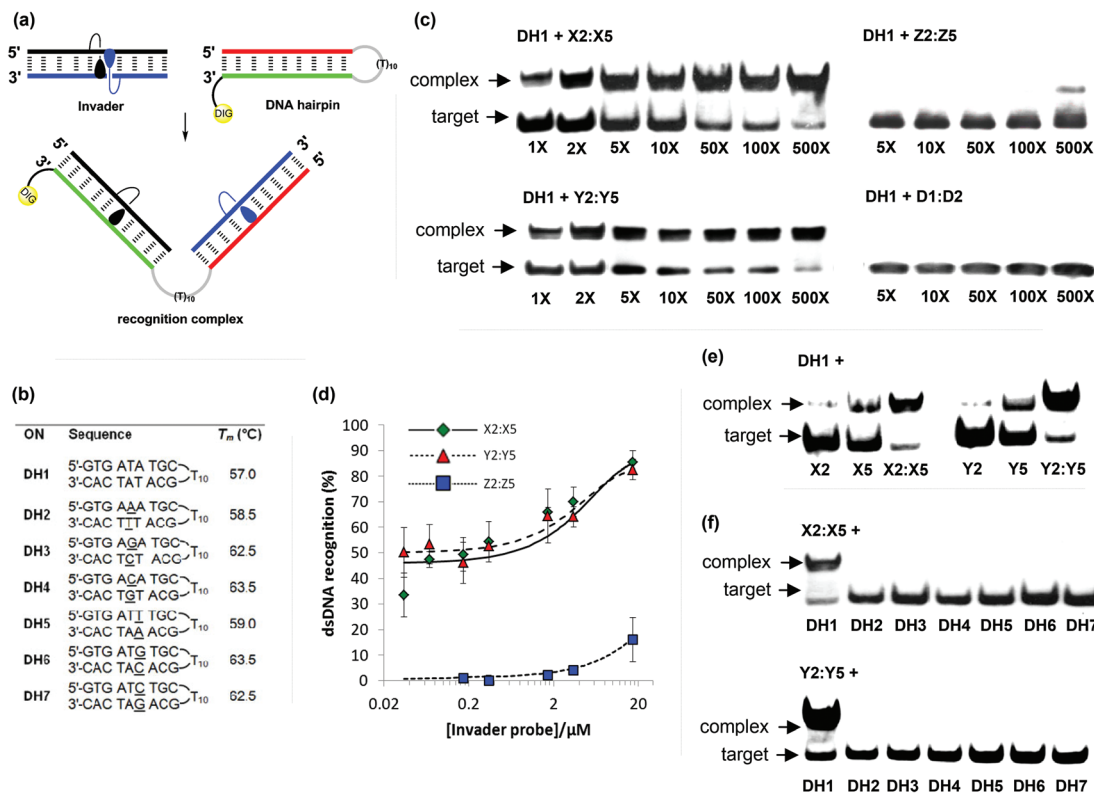


Fig. 6 Recognition of model dsDNA targets using activated probe duplexes. (a) Illustration of recognition process; (b) sequences and intramolecular T_m 's of DNA hairpins with isosequential (DH1) or mismatched stems (DH2–DH7) relative to B2:B5 probes (for conditions of thermal denaturation experiments, see Table 1); (c) representative gel electrophoretograms illustrating recognition of DH1 using 1- to 500-fold molar excess of X2:X5, Y2:Y5, Z2:Z5, or unmodified D1:D4; (d) dose-response curves (average of three independent experiments; error bars represent standard deviation); (e) incubation of DH1 with 500-fold molar excess of single-stranded X2, X5, Y2 or Y5 or double-stranded X2:X5 or Y2:Y5; (f) gel electrophoretograms illustrating incubation of DH1–DH7 with 500-fold molar excess of X2:X5 or Y2:Y5. Experimental conditions for electrophoretic mobility shift assay: separately pre-annealed targets (34.4 nM) and probes (variable concentration) were incubated for 15 h at ambient temperature in 1× HEPES buffer (50 mM HEPES, 100 mM NaCl, 5 mM MgCl₂, 10% sucrose, 1.4 mM spermine tetrahydrochloride, pH 7.2) and then run on 16% non-denaturing PAGE (performed at 70 V, 2 h, ~4 °C) using 0.5× TBE as a running buffer (45 mM Tris, 45 mM boric acid, 1 mM EDTA); DIG: digoxigenin.

nition efficiency of Invader probes, especially considering that detection of chromosomal DNA using Invaders based on the slightly less efficient 2'-O-(pyren-1-yl)-RNA monomers already has been demonstrated.¹³ Proof-of-concept studies aiming at expanding the repertoire of applications for Invader probes in molecular biology, nucleic acid diagnostics and medicinal chemistry are ongoing and will be reported in due course.

Experimental

Preparation of 2-pyrenemethanol

2-Pyrenemethanol was obtained from 4,5,9,10-tetrahydropyrene by combining known literature protocols.²² First, a solution of bromine (1.24 mL, 24.2 mmol) in anhydrous DMF (40 mL) was added dropwise over 2 h to a room temperature solution of 4,5,9,10-tetrahydropyrene^{22a} (5.00 g, 24.2 mmol) in anhydrous DMF (40 mL). The mixture was stirred for 4 h at rt after completed addition, poured into cold water and stirred overnight. The resulting precipitate was filtered, washed with water, dried and purified by silica gel column chromatography

(petroleum ether) to provide an off-white solid material (5.50 g), which was used in the next step without further purification.

The solid material was dissolved in anhydrous benzene (400 mL) along with DDQ (14.4 g, 63.6 mmol), and the solution was refluxed under an argon atmosphere for 4 h. The resulting slurry was filtered through a pad of celite, which was thoroughly washed with benzene. The filtrate was washed with aq. NaOH (10% v/v, 3 × 100 mL) and water (3 × 100 mL). The organic phase was evaporated to dryness and the resulting residue purified by silica gel column chromatography (petroleum ether) to afford a pale yellow solid material (4.60 g), which was used in the next step without further purification.

Next, *n*-butyllithium (9.82 mL, 24.5 mmol, 2.5 M in hexane) was added slowly to a −78 °C solution of the solid material in anhydrous ether–THF (350 mL, 1 : 1 v/v). The brightly red solution was stirred at −78 °C for 2 h and then slowly treated with anhydrous DMF (2.6 mL, 33.5 mmol). The reaction mixture was stirred at −30 to −50 °C for 1 h and then at rt for an additional 15 h. At this point, the reaction mixture was poured into ice-cold water (~100 mL) and the aqueous phase was



extracted with EtOAc (3 × 100 mL). The combined organic layers were evaporated to near dryness and the resulting residue was purified by silica gel column chromatography (0–3% CH₂Cl₂ in petroleum ether, v/v) to afford a brightly yellow solid material (3.50 g), which was used in the next step without further purification.

Lastly, sodium borohydride (0.69 g, 18.2 mmol) was added to a solution of this solid material in anhydrous THF (120 mL) and the reaction mixture was stirred overnight at rt. The reaction mixture was cooled on an ice bath and aqueous NaHCO₃ (10% v/v, 50 mL) was added carefully. The aqueous layer was extracted with EtOAc (3 × 100 mL) and the combined organic layers were evaporated to dryness. The resulting residue was purified by silica gel column chromatography (0–1% MeOH in CH₂Cl₂, v/v) to afford 2-pyrenemethanol^{22c} (3.45 g, 62% over four steps) as a white solid material. *R*_f = 0.5 (10% MeOH in CH₂Cl₂, v/v); MALDI-HRMS *m/z* 232.0875 ([M]⁺, C₁₇H₁₂O, Calc. 232.0888); ¹H NMR (DMSO-*d*₆): δ 8.28 (d, 2H, *J* = 7.8 Hz, H6/H8), 8.25 (s, 2H, H1/H3), 8.17 (s, 4H, H4/H5/H9/H10), 8.03–8.06 (ap t, 1H, *J* = 7.5 Hz, H7), 5.52 (t, 1H, ex, *J* = 5.7 Hz, OH), 4.97 (d, 2H, *J* = 5.7 Hz, CH₂OH); ¹³C NMR (DMSO-*d*₆) δ 140.7 (C2), 130.5, 130.4, 127.31 (C4/C5),³⁴ 127.29 (C9/C10), 125.9 (C7), 124.9 (C6/C8), 123.8, 123.0 (C1/C3), 122.9, 63.2 (CH₂Py).

Preparation of 4-pyrenemethanol

4-Pyrenemethanol was obtained from 4,5,9,10-tetrahydropyrene by combining known literature protocols²³ and protocols for the synthesis of 2-pyrenemethanol. First, a solution of bromine (0.62 mL, 12.0 mmol) in glacial acetic acid (25 mL) was added dropwise over 1 h to a room temperature solution of 4,5,9,10-tetrahydropyrene 3^{22a} (2.50 g, 12.0 mmol) in glacial acetic acid (25 mL). After ended addition, the reaction mixture was heated to 80 °C for 30 min, and then slowly cooled to rt, leading to the formation of a precipitate, which was isolated to provide a white solid material (2.90 g, 10.1 mmol) which used in the next step without further purification.

The solid material and DDQ (7.56 g, 33.3 mmol) were dissolved in anhydrous benzene (300 mL) and refluxed for 4 h. The resulting slurry was filtered through a celite pad, which was thoroughly washed with benzene. The filtrate was washed with aq. NaOH (10% v/v, 3 × 80 mL) and water (3 × 80 mL). The organic phase was evaporated to dryness and the resulting residue purified *via* silica gel column chromatography (petroleum ether) to obtain a pale yellow solid (2.00 g), which was used in the next step without further purification.

Next, *n*-butyllithium (4.27 mL, 10.7 mmol, 2.5 M in hexane) was added slowly to a –78 °C solution of this solid material in anhydrous ether–THF (170 mL, 1 : 1 v/v). The brightly red solution was stirred at –78 °C for 2 h and then treated slowly with anhydrous DMF (1.12 mL, 14.6 mmol). The reaction mixture was stirred at –30 to –50 °C for 1 h and then at rt for additional 15 h. The reaction mixture was poured into ice-cold water (~50 mL), the aqueous phase was extracted with EtOAc (3 × 50 mL), and the combined organic phase evaporated to dryness. The resulting residue was purified by silica gel

column chromatography (0–3% CH₂Cl₂ in pet ether, v/v) to afford a brightly yellow solid material (1.20 g), which was used in the next step without further purification.

Finally, NaBH₄ (0.24 g, 6.25 mmol) was added to a solution of the solid material in anhydrous THF (80 mL) and the reaction mixture was stirred overnight at rt. The reaction mixture was cooled on an ice-bath and aqueous NaHCO₃ (10% v/v, 20 mL) was added carefully. The aqueous layer was extracted with EtOAc (3 × 80 mL) and the combined organic layers were evaporated to dryness. The resulting residue was purified by silica gel column chromatography (0–1% MeOH in CH₂Cl₂, v/v) to afford 4-pyrenemethanol (1.15 g, 41% over four steps) as a white solid material. *R*_f = 0.5 (10% MeOH in CH₂Cl₂, v/v); MALDI-HRMS *m/z* 232.0881 ([M]⁺, C₁₇H₁₂O, Calc. 232.0888); ¹H NMR (DMSO-*d*₆): δ 8.39–8.41 (dd, 1H, *J* = 7.4 Hz, 0.8 Hz, H3), 8.26–8.32 (m, 3H, H1/H6/H8), 8.24 (s, 1H, H5), 8.17–8.21 (2d, 2H, *J* = 8.8 Hz, H9/H10), 8.05–8.11 (m, 2H, H2/H7), 5.53 (t, 1H, ex, *J* = 5.5 Hz, OH), 5.19–5.21 (dd, 2H, *J* = 5.5 Hz, 0.8 Hz, CH₂Py); ¹³C NMR (DMSO-*d*₆) δ 137.1, 130.9, 130.4, 130.3, 128.9, 127.4 (C9/C10), 127.1 (C9/C10), 126.2 (C2/C7), 125.9 (C2/C7), 125.1 (C1/C6/C8), 125.0 (C1/C6/C8), 124.8 (C1/C6/C8), 124.3 (C5), 124.0, 123.2, 121.2 (C3), 61.4 (CH₂Py).

General O2'-alkylation protocol for the preparation of 2Y/2Z (description for ~4.4 mmol scale)

The appropriate pyrenemethanol, solid NaHCO₃ and borane (1.0 M solution in THF) were placed in a pressure tube, suspended in anhydrous DMSO and stirred under an argon atmosphere at rt until effervescence ceased (~10 min). At this point, O2,O2'-anhydrouridine 1²⁰ was added (specific quantities of substrates and reagents are given below), the pressure tube was purged with argon and sealed, and the mixture was heated at ~140 °C until analytical TLC indicated full conversion (~3 h). After cooling to rt, the reaction mixture was poured into water (~50 mL), stirred for 30 min and diluted with EtOAc (~100 mL). The organic phase was washed with water (4 × 50 mL), evaporated to dryness and the resulting crude purified by silica gel column chromatography (2–4%, MeOH in CH₂Cl₂, v/v) to afford a residue, which was further purified through precipitation from refluxing methanol (30 min) to afford nucleoside 2 (yield specified below).

2'-O-(Pyren-2-yl)methyl-uridine (2Y)

O2,O2'-Anhydrouridine 1 (1.00 g, 4.42 mmol), 2-pyrenemethanol (2.05 g, 8.84 mmol), NaHCO₃ (74.3 mg, 0.88 mmol), BH₃ in THF (2.21 mL, 2.21 mmol) and anhydrous DMSO (8 mL) were mixed, reacted, worked up, and purified as described above to afford nucleoside 2Y (0.60 g, 30%) as a white solid material. *R*_f: 0.4 (10% MeOH in CH₂Cl₂, v/v); MALDI-HRMS *m/z* 481.1386 ([M + Na]⁺, C₂₆H₂₂N₂O₆·Na⁺, Calc. 481.1370); ¹H NMR (DMSO-*d*₆):³⁵ δ 11.38 (s, 1H, ex, NH), 8.28–8.30 (d, 2H, *J* = 7.8 Hz, H6_{Py}, H8_{Py}), 8.23 (s, 2H, H1_{Py}, H3_{Py}), 8.17–8.20 (d, 2H, *J* = 8.8 Hz, H4_{Py}, H10_{Py}), 8.10–8.13 (d, 2H, *J* = 8.8 Hz, H5_{Py}, H9_{Py}), 8.07 (t, 1H, *J* = 7.8 Hz, H7_{Py}), 7.85 (d, 1H, *J* = 8.0 Hz, H6), 6.09 (d, 1H, *J* = 4.9 Hz, H1'), 5.51 (d, 1H, *J* = 8.0 Hz, H5), 5.35 (d, 1H, ex, *J* = 5.7 Hz, 3'-OH), 5.16–5.19 (d, 1H, *J* = 12.7 Hz,



CH₂Py), 5.11 (t, 1H, ex, J = 4.9 Hz, 5'-OH), 5.03–5.06 (d, 1H, J = 12.7 Hz, CH₂Py), 4.23–4.26 (m, 1H, H3'), 4.12–4.15 (m, 1H, H2'), 3.98–4.00 (m, 1H, H4'), 3.65–3.69 (m, 1H, H5'), 3.57–3.62 (m, 1H, H5'); ¹³C NMR (DMSO-*d*₆): δ 163.0, 150.6, 140.2 (C6), 136.3, 130.54, 130.50, 127.5 (C4_{Py}, C10_{Py}), 127.2 (C5_{Py}, C9_{Py}), 126.1 (C7_{Py}), 125.1 (C6_{Py}, C8_{Py}), 123.7 (C1_{Py}, C3_{Py}), 123.2, 101.8 (C5), 86.4 (C1'), 85.2 (C4'), 80.8 (C2'), 71.3 (CH₂Py), 68.4 (C3'), 60.5 (C5').

2'-O-(Pyren-4-yl)methyl-uridine (2Z)

O₂O₂'-Anhydrouridine **1** (0.50 g, 2.21 mmol), 4-pyrenemethanol (1.03 g, 4.42 mmol), NaHCO₃ (37.1 mg, 0.44 mmol), BH₃ in THF (1.10 mL, 1.10 mmol) and anhydrous DMSO (5 mL) were mixed, reacted, worked up, and purified as described above to afford nucleoside **2Y** (200 mg, 20%) as a white solid. *R*_f: 0.4 (10% MeOH in CH₂Cl₂, v/v); MALDI-HRMS *m/z* 481.1399 ([M + Na]⁺, C₂₆H₂₂N₂O₆·Na⁺, Calc. 481.1370; ¹H NMR (DMSO-*d*₆): δ 11.28 (s, 1H, ex, NH), 8.42 (d, 1H, J = 7.8 Hz, Py), 8.29–8.31 (m, 2H, Py), 8.24–8.26 (d, 1H, J = 7.3 Hz, Py), 8.18–8.22 (m, 3H, Py), 8.07 (t, 1H, J = 7.5 Hz, Py), 8.03 (dd, 1H, J = 7.8 Hz, 7.5 Hz, Py), 7.77 (d, 1H, J = 8.0 Hz, H6), 6.08 (d, 1H, J = 5.4 Hz, H1'), 5.39–5.43 (m, 3H, 1 ex, 3'-OH, H5 and CH₂Py), 5.21–5.24 (d, 1H, J = 12.5 Hz, CH₂Py), 5.09 (t, 1H, ex, 5'-OH), 4.29–4.33 (m, 1H, H3'), 4.20–4.23 (m, 1H, H2'), 3.97–4.00 (m, 1H, H4'), 3.63–3.68 (m, 1H, H5'), 3.57–3.62 (m, 1H, H5'); ¹³C NMR (DMSO-*d*₆): δ 162.8, 150.5, 140.0 (C6), 132.8, 130.8, 130.5, 129.9, 129.0, 127.5 (Py), 127.0 (Py), 126.9 (Py), 126.3 (Py), 125.9 (Py), 125.3 (Py), 125.2 (Py), 124.1, 123.5, 121.7 (Py), 101.6 (C5), 86.0 (C1'), 85.4 (C4'), 80.7 (C2'), 70.2 (CH₂Py), 68.4 (C3'), 60.6 (C5').

General O5'-DMTr-protection protocol for the preparation of 3Y/3Z (description for ~1 mmol scale)

The appropriate nucleoside **2** (specific quantities given below) was co-evaporated twice with anhydrous pyridine and redissolved in anhydrous pyridine. To this was added 4,4'-dimethoxytritylchloride (DMTrCl) and catalytic *N,N*-dimethyl-4-aminopyridine (DMAP), and the reaction mixture was stirred at rt under an argon atmosphere until TLC indicated complete conversion (~14 h). The reaction mixture was diluted with CH₂Cl₂ (40 mL) and the organic phase was sequentially washed with water (2 × 30 mL) and sat. aq. NaHCO₃ (2 × 50 mL). The organic phase was evaporated to near dryness and the resulting crude co-evaporated with abs. EtOH and toluene (2 : 1, v/v, 3 × 3 mL) and purified by silica gel column chromatography (0–5%, MeOH in CH₂Cl₂, v/v) to afford nucleoside **3** (yield specified below).

5'-O-(4,4'-Dimethoxytrityl)-2'-O-(pyren-2-yl)methyl-uridine (3Y)

Nucleoside **2Y** (500 mg, 1.09 mmol), DMTrCl (0.64 g, 1.63 mmol) and DMAP (~9 mg) in anhydrous pyridine (10 mL) were mixed, reacted, worked up and purified as described above to afford **3Y** (0.57 g, 68%) as a pale yellow foam. *R*_f: 0.6 (5% MeOH in CH₂Cl₂, v/v); MALDI-HRMS *m/z* 783.2706 ([M + Na]⁺, C₄₇H₄₀N₂O₈·Na⁺, Calc. 783.2677); ¹H NMR (DMSO-*d*₆): δ 11.43 (s, 1H, ex, NH), 8.27–8.31 (m, 4H, H1_{Py},

H3_{Py}, H6_{Py}, H8_{Py}), 8.17–8.20 (d, 2H, J = 8.9 Hz, H4_{Py}, H10_{Py}), 8.11–8.13 (d, 2H, J = 8.9 Hz, H5_{Py}, H9_{Py}), 8.07 (t, 1H, J = 7.5 Hz, H7_{Py}), 7.67 (d, 1H, J = 8.0 Hz, H6), 7.31–7.33 (m, 2H, DMTr), 7.22–7.26 (m, 2H, DMTr), 7.15–7.20 (m, 5H, DMTr), 6.78–6.84 (m, 4H, DMTr), 6.07–6.08 (d, 1H, J = 3.9 Hz, H1'), 5.45 (d, 1H, ex, J = 6.7 Hz, 3'-OH), 5.13–5.23 (m, 3H, H5, CH₂Py), 4.31–4.35 (m, 1H, H3'), 4.18–4.20 (m, 1H, H2'), 4.09–4.12 (m, 1H, H4'), 3.68 (s, 3H, CH₃O), 3.65 (s, 3H, CH₃O), 3.30–3.34 (m, 1H, H5'; overlap with H₂O), 3.23–3.26 (m, 1H, H5'); ¹³C NMR (DMSO-*d*₆): δ 162.9, 158.1, 158.0, 150.4, 144.4, 140.1 (C6), 136.2, 135.3, 135.0, 130.6, 130.5, 129.7 (DMTr), 129.6 (DMTr), 127.8 (DMTr), 127.6 (DMTr), 127.5 (Py), 127.2 (Py), 126.7 (Py), 126.1 (Py), 125.1 (Py), 123.68, 123.66 (Py), 123.2, 113.2 (DMTr), 113.1 (DMTr), 101.5 (C5), 87.2 (C1'), 85.9, 82.9 (C4'), 80.6 (C2'), 71.3 (CH₂Py), 68.7 (C3'), 62.7 (C5'), 54.94 (CH₃O), 54.92 (CH₃O). A minor impurity of chloroform at 79.1 ppm was identified in the ¹³C NMR.

5'-O-(4,4'-Dimethoxytrityl)-2'-O-(pyren-4-yl)methyl-uridine (3Z)

Nucleoside **2Z** (150 mg, 0.33 mmol), DMTrCl (191 mg, 0.49 mmol) and DMAP (~4 mg) in anhydrous pyridine (5 mL) were mixed, reacted, worked up and purified as described above to afford **3Z** (200 mg, 80%) as a pale yellow foam. *R*_f: 0.6 (5% MeOH in CH₂Cl₂, v/v); MALDI-HRMS *m/z* 783.2695 ([M + Na]⁺, C₄₇H₄₀N₂O₈·Na⁺, Calc. 783.2677); ¹H NMR (DMSO-*d*₆): δ 11.36 (s, 1H, ex, NH), 8.45 (d, 1H, J = 7.8 Hz, Py), 8.17–8.33 (m, 6H, Py), 8.05 (t, 2H, H2_{Py}, H7_{Py}), 7.60 (d, 1H, J = 8.2 Hz, H6), 7.30–7.32 (m, 2H, DMTr), 7.17–7.25 (m, 7H, DMTr), 6.78–6.84 (m, 4H, DMTr), 6.07 (d, 1H, J = 4.1 Hz, H1'), 5.49 (d, 1H, J = 6.2 Hz, ex, 3'-OH), 5.44–5.46 (m, 1H, J = 12.7 Hz, CH₂Py), 5.30–5.34 (d, 1H, J = 12.7 Hz, CH₂Py), 5.08 (d, 1H, J = 8.2 Hz, H5), 4.38–4.42 (m, 1H, H3'), 4.26–4.28 (m, 1H, H2'), 4.09–4.12 (m, 1H, H4'), 3.71 (s, 3H, CH₃O), 3.69 (s, 3H, CH₃O), 3.30–3.34 (m, 1H, H5' – partial overlap with H₂O), 3.22–3.25 (m, 1H, H5'); ¹³C NMR (DMSO-*d*₆): δ 162.8, 158.1, 158.0, 150.4, 144.4, 140.0 (C6), 135.3, 135.0, 132.7, 130.9, 130.5, 129.9, 129.7 (DMTr), 129.6 (DMTr), 129.0, 127.8 (DMTr), 127.6 (DMTr), 127.5 (Py), 127.1 (Py), 126.8 (Py), 126.7 (DMTr), 126.3 (C2_{Py}), 126.0 (C7_{Py}), 125.29 (Py), 125.25 (Py), 124.1, 123.5, 121.7 (Py), 113.2 (DMTr), 113.1 (DMTr), 101.4 (C5), 86.9 (C1'), 85.9, 83.2 (C4'), 80.5 (C2'), 70.2 (CH₂Py), 68.7 (C3'), 62.8 (C5'), 55.0 (CH₃O).

General O3'-phosphitylation protocol for the preparation of 4Y and 4Z (description for ~0.25 mmol scale)

The appropriate nucleoside **3** (specific quantities given below) was co-evaporated with anhydrous 1,2-dichloroethane and redissolved in anhydrous CH₂Cl₂. To this was added *N,N*-diisopropylethylamine (DIPEA) and 2-cyanoethyl-*N,N*-diisopropylchlorophosphoramidite (PCL-reagent) and the reaction mixture was stirred at rt under an argon atmosphere until TLC indicated complete conversion (~3 h), whereupon abs. EtOH (0.3 mL) and CH₂Cl₂ (5 mL) were sequentially added. The organic phase was washed with sat. aq. NaHCO₃ (2 mL), evaporated to near dryness, and the resulting residue purified by silica gel column chromatography (40–70% EtOAc in



petroleum ether, v/v) to afford phosphoramidite **4** (yield specified below).

3'-O-(2-Cyanoethoxy(diisopropylamino)phosphinyl)-5'-O-(4,4'-dimethoxytrityl)-2'-O-(pyren-2-yl)methyl-uridine (4Y**)**

Nucleoside **3Y** (145 mg, 0.19 mmol), DIPEA (136 μ L, 0.76 mmol) and PCI-reagent (85 μ L, 0.38 mmol) in anhydrous CH_2Cl_2 (5 mL) were mixed, reacted, worked up and purified as described above to afford phosphoramidite **4Y** (170 mg, 93%) as a white foam. R_f : 0.8 (5% MeOH in CH_2Cl_2 , v/v); MALDI-HRMS m/z 983.3746 ($[\text{M} + \text{Na}]^+$, $\text{C}_{56}\text{H}_{57}\text{N}_4\text{O}_9\text{P}\cdot\text{Na}^+$, Calc. 983.3755); ^{31}P NMR (CDCl_3): δ 150.5, 150.2.

3'-O-(2-Cyanoethoxy(diisopropylamino)phosphinyl)-5'-O-(4,4'-dimethoxytrityl)-2'-O-(pyren-4-yl)methyl-uridine (4Z**)**

Nucleoside **3Z** (190 mg, 0.25 mmol), DIPEA (178 μ L, 0.99 mmol) and PCI-reagent (111 μ L, 0.49 mmol) in anhydrous CH_2Cl_2 (5 mL) were mixed, reacted, worked up and purified as described above to afford phosphoramidite **4Z** (220 mg, 92%) as a white foam. R_f : 0.8 (5% MeOH in CH_2Cl_2 , v/v); MALDI-HRMS m/z 983.3766 ($[\text{M} + \text{Na}]^+$, $\text{C}_{56}\text{H}_{57}\text{N}_4\text{O}_9\text{P}\cdot\text{Na}^+$, Calc. 983.3755); ^{31}P NMR (CDCl_3) δ 150.1, 150.0.

Protocol – synthesis and purification of ONs

Synthesis of **Y/Z**-modified ONs was performed on an automated DNA synthesizer using 0.2 μ mol scale succinyl linked LCAA-CPG (long chain alkyl amine controlled pore glass) columns with a pore size of 500 Å. Standard protocols for incorporation of DNA phosphoramidites were used. A ~50-fold molar excess of modified phosphoramidites in anhydrous acetonitrile (at 0.05 M), along with extended oxidation (45 s) and coupling times (activator: 0.01 M 4,5-dicyanoimidazole, 15 min) was used during hand-couplings, which resulted in stepwise coupling yields of **4Y** and **4Z** of ~99% and ~98%, respectively. Cleavage from solid support and removal of protecting groups was accomplished upon treatment with 32% aq. ammonia (55 °C, 24 h). The crude ONs were purified *via* ion-pair reverse phase HPLC (XTerra MS C18 column) using 0.05 M triethylammonium acetate–water/acetonitrile gradient, followed by detritylation (80% aq. AcOH, 20 min), and precipitation (NaOAc–NaClO₄–acetone, –18 °C for 12–16 h). The identity of synthesized ONs was established through MALDI-MS analysis (Table S1†) recorded in positive ions mode on a Quadrupole Time-Of-Flight Tandem Mass Spectrometer (Q-TOF Premiere) equipped with a MALDI source (Waters Micromass LTD, U.K) using anthranilic acid as a matrix, while purity (>85%) was verified by ion-pair reverse phase HPLC running in analytical mode.

Protocol – thermal denaturation studies

Concentrations of ONs were estimated using the following extinction coefficients for DNA (OD μmol^{-1}): G (12.01), A (15.20), T (8.40), C (7.05); for RNA (OD μmol^{-1}): G (13.70), A (15.40), U (10.00), C (9.00); for pyrene (OD μmol^{-1}): (22.4)³¹ Strands were thoroughly mixed and denatured by heating to 70–85 °C, followed by cooling to the starting temperature of

the experiment. Quartz optical cells with a path length of 1.0 cm were used. Thermal denaturation temperatures (T_m 's) of duplexes (1.0 μM final concentration of each strand) were measured on a Cary 100 UV/VIS spectrophotometer equipped with a 12-cell Peltier temperature controller and determined as the maximum of the first derivative of the thermal denaturation curve (A_{260} vs. T) recorded in medium salt buffer (T_m buffer: 100 mM NaCl, 0.1 mM EDTA, and pH 7.0 adjusted with 10 mM Na₂HPO₄ and 5 mM Na₂HPO₄). The temperature of the denaturation experiments ranged from at least 15 °C below T_m to 20 °C above T_m (although not below 3 °C). A temperature ramp of 0.5 °C min^{–1} was used in all experiments. Reported T_m -values are averages of two experiments within ± 1.0 °C.

Protocol – determination of thermodynamic parameters

Thermodynamic parameters for duplex formation were determined through baseline fitting of denaturation curves (van't Hoff analysis) using software provided with the UV/VIS spectrometer. Bimolecular reactions, two-state melting behavior, and a heat capacity change of $\Delta C_p = 0$ upon hybridization were assumed.³⁰ A minimum of two experimental denaturation curves were each analyzed at least three times to minimize errors arising from baseline choice. Averages and standard deviations are listed.

Protocol – absorption spectra

UV-vis absorption spectra (range: 300–400 nm) were recorded at 5 °C using the same samples and instrumentation as in the thermal denaturation experiments.

Protocol – steady-state fluorescence emission spectra

Steady-state fluorescence emission spectra of ONs modified with pyrene-functionalized monomers **X–Z** and the corresponding duplexes with complementary DNA/RNA targets, were recorded in non-deoxygenated thermal denaturation buffer (each strand at 1.0 μM concentration) and obtained as an average of five scans using an excitation wavelength of $\lambda_{\text{ex}} = 350, 345$ or 340 nm for **X**-, **Y**- or **Z**-modified ONs, excitation slit 5.0 nm, emission slit 2.5 nm and a scan speed of 600 nm min^{–1}. Experiments were determined at 5 °C to ascertain maximal hybridization of probes to DNA/RNA targets (a stream of nitrogen was blown in to the chamber to prevent condensation).

Protocol – molecular modeling

The initial structures of the **X/Y/Z**-modified DNA duplexes were generated by building and modifying a standard *B*-type DNA duplex using MacroModel v9.8.³⁷ The charge of the phosphodiester backbone was neutralized with sodium ions, which were placed 3.0 Å from non-bridging oxygen atoms and restrained to this distance by a force constant of 100 kJ mol^{–1} Å^{–2}. A preliminary minimization was carried out using the Polak–Ribiere conjugate gradient method (convergence criteria 0.1 kJ mol^{–1} Å^{–1}), the AMBER94 force field³⁸ with the improved parambsc0 parameter set,³⁹ and the implicit GB/SA continuum solvation model⁴⁰ as implemented in MacroModel



v9.8; all atoms were frozen except those in the O2'-substituent of the modified monomers. Non-bonded interactions were treated with extended cut-offs (van der Waals 8.0 Å and electrostatics 20.0 Å).

The seed structure for **X2:D4** was generated by taking previously reported NMR data on X-modified DNA duplexes into account.³¹ Thus, distances between atoms for which NOE contacts have been reported were constrained to 3.0–5.0 Å by a force constant of 100 kJ mol⁻¹ Å⁻² (*i.e.*, distances between (i) pyrene and H5 of the modified uridine, (ii) pyrene and H2/H8 of the 3'-flanking adenosine, and (iii) pyrene and H6 of the thymine opposite of the 3'-flanking adenosine). The structure was minimized as described above with the following considerations: (i) all atoms were allowed to move freely during minimization, (ii) sodium ions were restrained as above, and (iii) hydrogen bonding of the outermost base pairs were restrained by a force constant of 100 kJ mol⁻¹ Å⁻² ((C)2–O...HN–2(G), distance 1.99 Å, (C)N3...H–N1(G), distance 1.90 Å and (C)4–NH...O–6(G), distance 1.69 Å). The resulting lowest energy structure formed the basis for construction of the seed structures for all other studied duplexes. Pyrene moieties of the other duplexes were manually adjusted to intercalated positions and the duplexes minimized as described for **X2:D4**. Seed structures were then submitted to 5 ns of stochastic dynamics (simulation temperature 300 K, time step 2.2 fs, SHAKE all bonds to hydrogen) using the same constraints as employed above (except for NOE constraints), during which 1000 structures were sampled and subsequently minimized. The resulting structures were analyzed using MacroModel v9.8 and the 3DNA web server.⁴¹

Protocol – electrophoretic mobility shift assay

This assay was performed, essentially as previously described.¹³ Unmodified DNA hairpins **DH1–DH7** were obtained from commercial sources and used without further purification. The DNA hairpins were 3'-DIG-labeled using the 2nd generation DIG Gel Shift Kit (Roche Applied Bioscience) as recommended by the manufacturer. DIG-labeled ONs obtained in this manner, were diluted and used without further purification in the recognition experiments. Pre-annealed probes (90 °C for 10 min, cooled to room temperature over 15 min) and DIG-labeled DNA hairpins (34.4 nM) were mixed and incubated in HEPES buffer (50 mM HEPES, 100 mM NaCl, 5 mM MgCl₂, 10% sucrose, 1.44 mM spermine tetrahydrochloride, pH 7.2) for 15 h at ambient temperature (~21 ± 3 °C). The reaction mixtures were then diluted with 6x DNA loading dye (Fermentas) and loaded onto a 16% non-denaturing polyacrylamide gel. Electrophoresis was performed using a constant voltage of 70 V for 2 h at ~4 °C. Gels were blotted onto a positively charged nylon membrane (Roche Applied Bioscience) using constant voltage with external cooling (100 V, ~4 °C). The membranes were exposed to anti-digoxigenin-AP F_{ab} fragments as recommended by the manufacturer of the DIG Gel Shift Kit, transferred to a hybridization jacket, and incubated with the substrate (CSPD) in detection buffer for 10 min at 37 °C. The chemiluminescence of the formed product was captured

on X-ray film, which was developed using an X-Omatic 1000A X-ray film developer (Kodak). The resulting bands were quantified using a Fluor-S MultiImager (Bio-Rad, Hercules, CA) equipped with Quantity One software. Invasion efficiency was determined as the intensity ratio between the recognition complex band and the total lane. An average of three independent experiments is reported along with standard deviations.

Definition of zipper nomenclature

The following nomenclature describes the relative arrangement between two monomers positioned on opposing strands in a duplex. The number *n* describes the distance measured in number of base pairs and has a positive value if a monomer is shifted toward the 5'-side of its own strand relative to a second reference monomer on the other strand. Conversely, *n* has a negative value if a monomer is shifted toward the 3'-side of its own strand relative to a second reference monomer on the other strand.

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- 35 The assignments of H4_{Py}/H10_{Py} and H5_{Py}/H9_{Py} (and the corresponding ^{13}C NMR signals) are interchangeable.
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Recognition of double-stranded DNA using energetically activated duplexes with interstrand zippers of 1-, 2- or 4-pyrenyl-functionalized O2'-alkylated RNA monomers

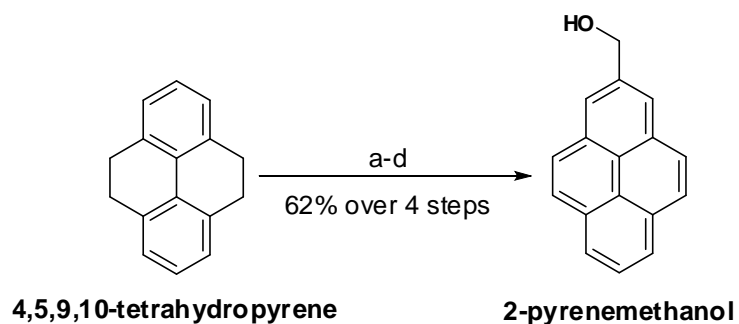
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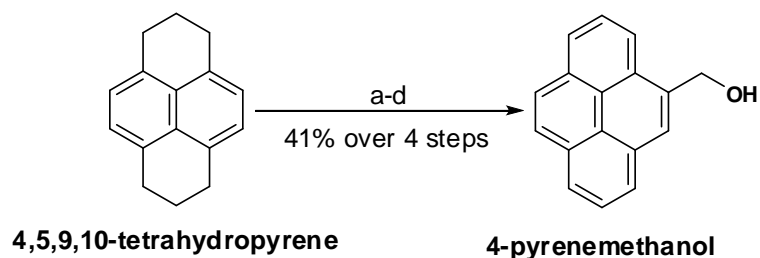
ELECTRONIC SUPPLEMENTARY INFORMATION

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General experimental section. Reagents and solvents were commercially available, of analytical grade and used without further purification. Petroleum ether of the distillation range 60-80 °C was used. Solvents were dried over activated molecular sieves: THF (3Å); CH₂Cl₂, 1,2-dichloroethane, *N,N'*-diisopropylethylamine and anhydrous DMSO (4Å). Anhydrous diethyl ether, DMF and benzene were obtained from commercial suppliers (<10 ppm water). Water content of anhydrous solvents was verified on Karl-Fisher apparatus. Reactions were conducted under argon whenever anhydrous solvents were used. Reactions were monitored by TLC using silica gel coated plates with a fluorescence indicator (SiO₂-60, F-254) which were visualized a) under UV light and/or b) by dipping in 5% conc. H₂SO₄ in absolute ethanol (v/v) followed by heating. Silica gel column chromatography was performed with Silica gel 60 (particle size 0.040–0.063 mm) using moderate pressure (pressure ball). Evaporation of solvents was carried out under reduced pressure at temperatures below 45 °C. After column chromatography, appropriate fractions were pooled, evaporated and dried at high vacuum for at least 12h to give the obtained products in high purity (>95%) as ascertained by 1D NMR techniques. Chemical shifts of ¹H NMR (500 MHz), ¹³C NMR (125.6 MHz), and/or ³¹P NMR (121.5 MHz) are reported relative to deuterated solvent or other internal standards (80% phosphoric acid for ³¹P NMR). Exchangeable (ex) protons were detected by disappearance of signals upon D₂O addition. Assignments of NMR spectra are based on 2D spectra (COSY, HSQC) and DEPT-spectra. Quaternary carbons are not assigned in ¹³C NMR but verified from HSQC and DEPT spectra (absence of signals). MALDI-HRMS spectra of compounds were recorded on a Waters Q-TOF Premiere mass spectrometer using 2,5-dihydroxybenzoic acid as a matrix and polyethylene glycol (PEG 600) as an internal calibration standard.



Scheme S1. Synthesis of 2-pyrenemethanol. Reagents and conditions: (a) Br₂, DMF, rt, 6h; (b) DDQ, benzene, reflux, 4h; (c) (i) *n*-BuLi, THF/ether, -78 °C for 2h; (ii) DMF, -30 to -50 °C for 1h, then rt for 15h; (d) NaBH₄, THF, rt, overnight.



Scheme S2. Synthesis of 4-pyrenemethanol. Reagents and conditions: (a) Br₂, AcOH, rt, 1h; (b) DDQ, benzene, reflux, 4h; (c) (i) *n*-BuLi, THF/ether, -78 °C for 2h; (ii) DMF, -50 to -30 °C for 1h, then rt for 15h; (d) NaBH₄, THF, rt, overnight.

Table S1. MALDI-MS of ONs modified with monomers **Y** and **Z**.^a

ONs	Sequence	Calc. m/z [M+H]	Found m/z [M+H]
Y1	5'-G <u>Y</u> G ATA TGC	2969	2969
Y2	5'-GTG A <u>Y</u> A TGC	2969	2969
Y3	5'-GTG ATA <u>Y</u> GC	2969	2969
Y4	3'-CAC <u>Y</u> AT ACG	2898	2898
Y5	3'-CAC TA <u>Y</u> ACG	2898	2898
Y6	3'-CAC <u>Y</u> A <u>Y</u> ACG	3114	3114
Z1	5'-G <u>Z</u> G ATA TGC	2969	2969
Z2	5'-GTG A <u>Z</u> A TGC	2969	2969
Z3	5'-GTG ATA <u>Z</u> GC	2969	2969
Z4	3'-CAC <u>Z</u> AT ACG	2898	2898
Z5	3'-CAC TA <u>Z</u> ACG	2898	2898
Z6	3'-CAC <u>Z</u> A <u>Z</u> ACG	3114	3114

^a For structures of monomers **Y** and **Z**, see Figure 1 in the main manuscript.

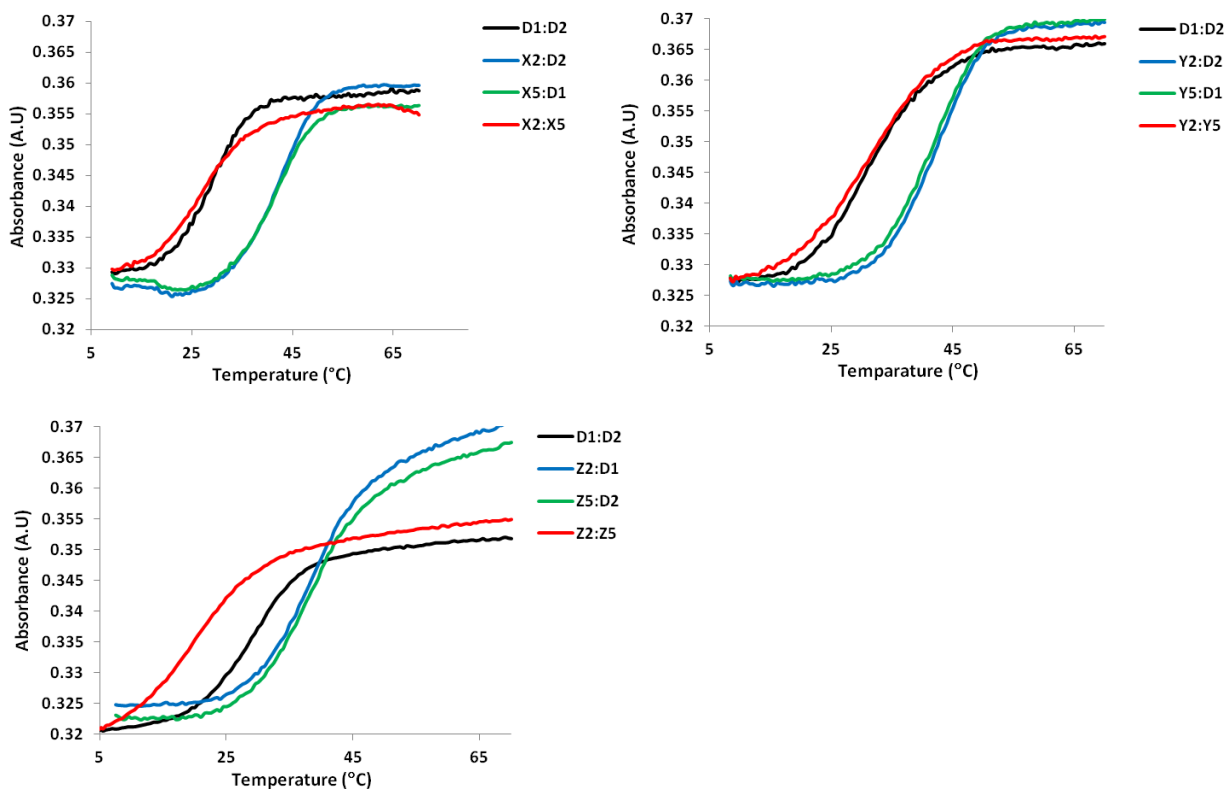


Figure S1. Representative thermal denaturation curves of **X/Y/Z**-modified duplexes and reference duplex **D1:D2**. For experimental conditions, see Table 1.

Table S2. DNA selectivity of **B1-B6**.^a

ON	Duplex	<u>B</u> =	$\Delta\Delta T_m/\text{mod (DNA-RNA)} [^{\circ}\text{C}]$		
			X	Y	Z
B1	5'-G <u>B</u> G ATA TGC		+7.0	+7.5	+6.0
B2	5'-GTG A <u>B</u> A TGC		+9.0	+6.5	+7.0
B3	5'-GTG ATA <u>B</u> GC		+8.0	+8.5	+9.0
B4	3'-CAC <u>B</u> AT ACG		+8.0	+7.5	+5.0
B5	3'-CAC TAB <u>B</u> ACG		+10.5	+6.5	+7.0
B6	3'-CAC <u>B</u> AB ACG		+7.3	+6.0	+5.5

^a DNA selectivity defined as $\Delta\Delta T_m/\text{mod (DNA-RNA)} = \Delta T_m/\text{mod (vs cDNA)} - \Delta T_m/\text{mod (vs cRNA)}$.

Additional discussion of binding specificity. **X2/Y2/Z2** display less efficient discrimination of mismatched RNA targets compared to reference strand **D1**, except when a mismatched G is opposite of the pyrene-functionalized monomers (Table S3).

ONs with two modifications positioned as next-nearest neighbors (**B6**-series) very efficiently discriminate DNA targets with a single mismatched nucleotide opposite of the central adenosine residue (Table S4). Similar observations have been observed with analogous ONs modified with other O2'-pyrene-functionalized RNA monomers or N2'-pyrene-functionalized 2'-*N*-methyl-2'-amino-DNA,^{S1} which suggests that this probe architecture may be a general strategy toward improving binding specificity. However, this hypothesis must be evaluated across different sequence contexts.

Table S3. Discrimination of mismatched RNA targets by **X2/Y2/Z2** and reference ONs.^a

ON	Sequence	B =	RNA: 3'-CAC U <u>B</u> U ACG			
			$T_m/^\circ\text{C}$		$\Delta T_m/^\circ\text{C}$	
			A	C	G	U
D1	5'-GTG ATA TGC		26.5	<-16.5	-4.5	<-16.5
X2 ^b	5'-GTG A <u>X</u> A TGC		31.0	-17.5	-3.5	-9.5
Y2	5'-GTG A <u>Y</u> A TGC		34.0	-13.5	-8.5	-13.5
Z2	5'-GTG A <u>Z</u> A TGC		27.5	-13.0	-7.0	-8.0

^a For conditions of thermal denaturation experiments, see Table 1. T_m 's of fully matched duplexes are shown in bold. ΔT_m = change in T_m relative to fully matched DNA:RNA duplex.

^b Data previously reported in reference S1. Included to facilitate direct comparison.

Table S4. Discrimination of mismatched DNA targets by **X6/Y6/Z6** and reference ONs.^a

ON	Sequence	B =	DNA : 5'-GTG A <u>B</u> A ACG			
			$T_m [^\circ\text{C}]$	$\Delta T_m [^\circ\text{C}]$		
			T	A	C	G
D2	3'-CAC TAT ACG		29.5	-17.0	-15.5	-9.0
X6	3'-CAC <u>X</u> A <u>X</u> ACG		43.5	-24.0	-17.0	-14.0
Y6	3'-CAC <u>Y</u> A <u>Y</u> ACG		45.5	-24.0	-16.0	-14.0
Z6	3'-CAC <u>Z</u> A <u>Z</u> ACG		32.5	-21.0	-16.0	-12.0

^a For conditions of thermal denaturation experiments, see Table 1. T_m 's of fully matched duplexes are shown in bold. ΔT_m = change in T_m relative to fully matched DNA:DNA duplex.

^b Data previously reported in reference S1. Included to facilitate direct comparison.

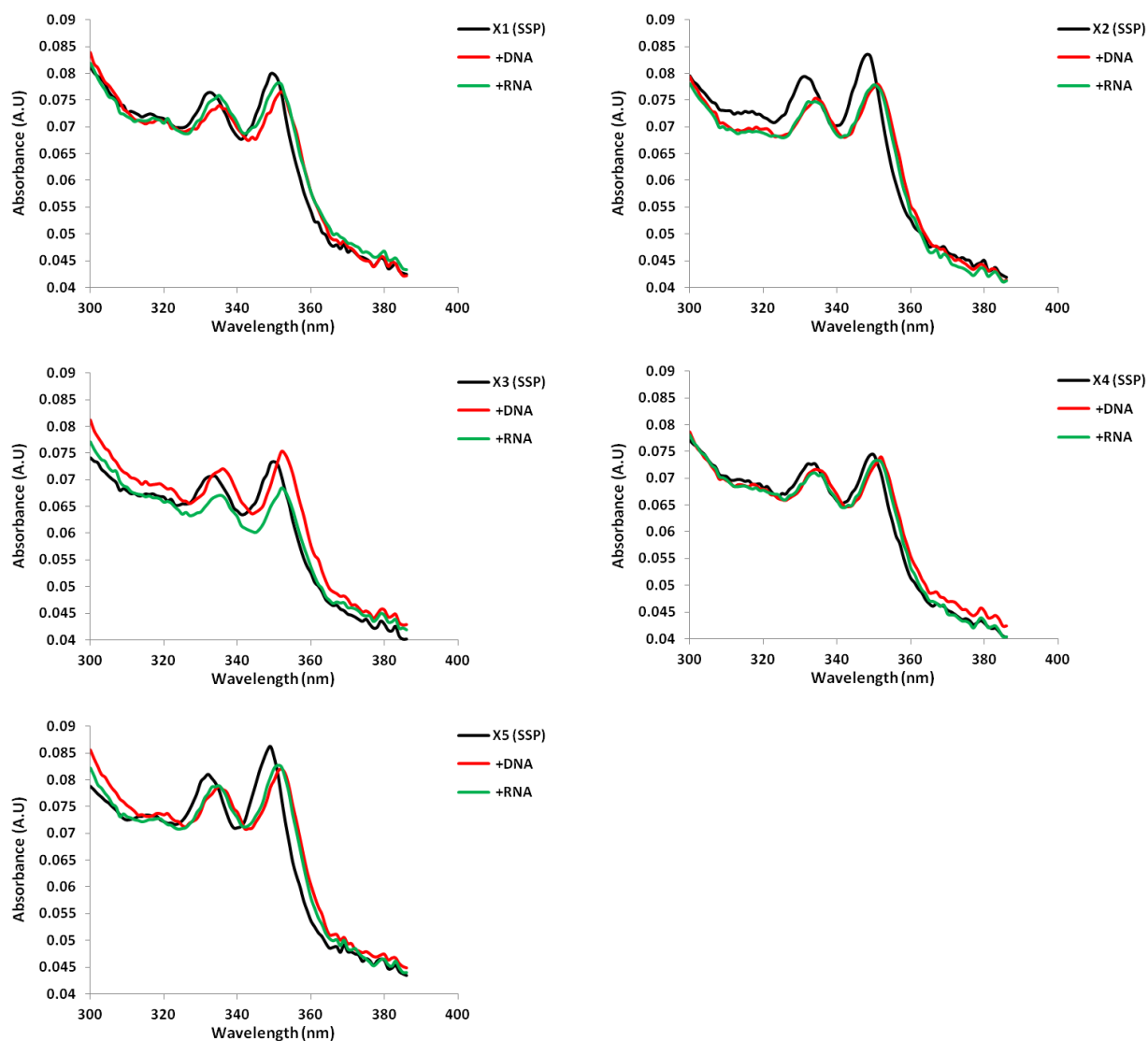


Figure S2. Absorbance spectra of single-stranded **X1-X5** and their corresponding duplexes with DNA/RNA targets. Spectra were recorded at $T = 5\text{ }^{\circ}\text{C}$ using each strand at $1.0\text{ }\mu\text{M}$ concentration in T_m buffer.

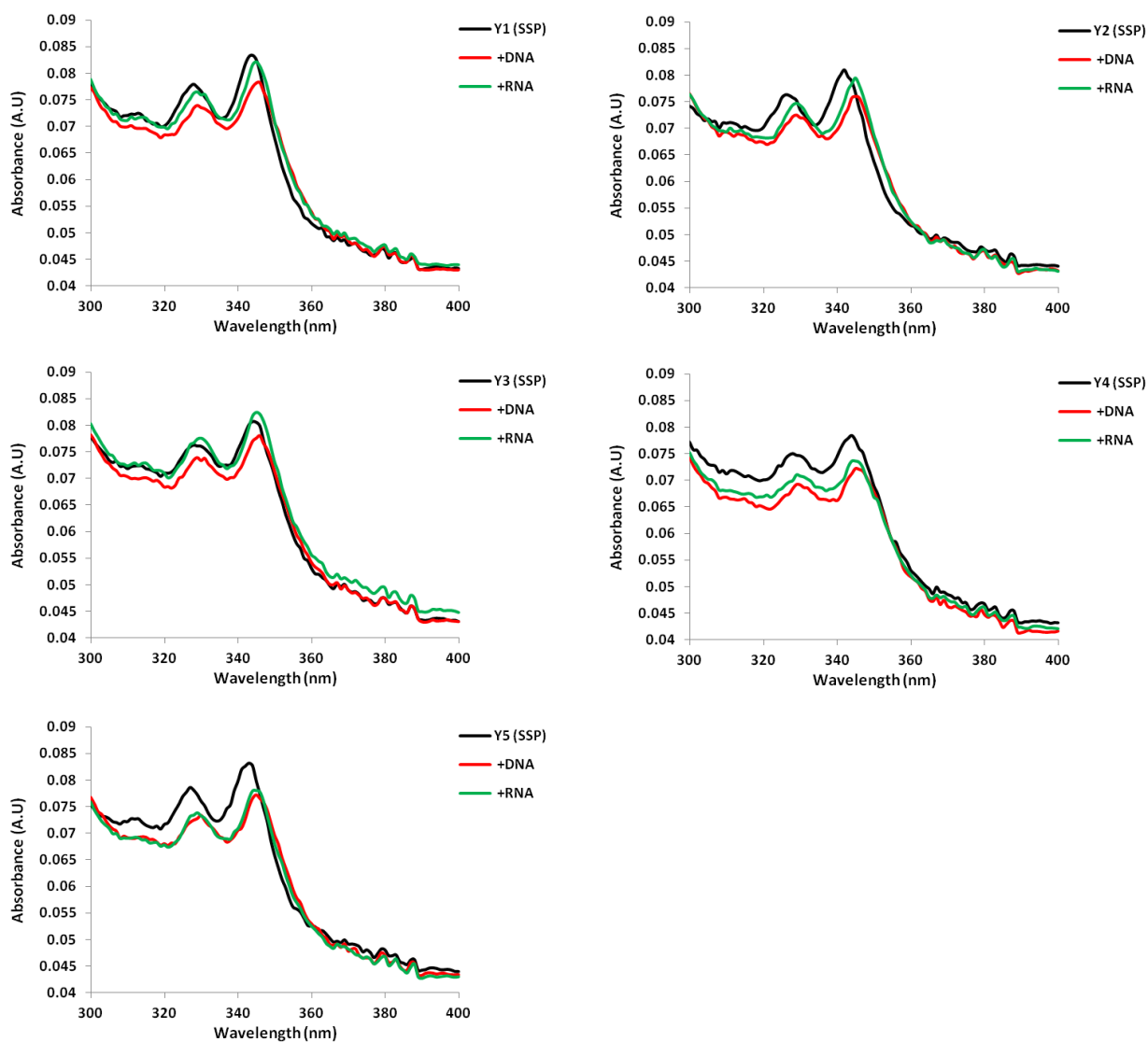


Figure S3. Absorbance spectra of single-stranded **Y1-Y5** and their corresponding duplexes with DNA/RNA targets. For conditions, see Fig. S2.

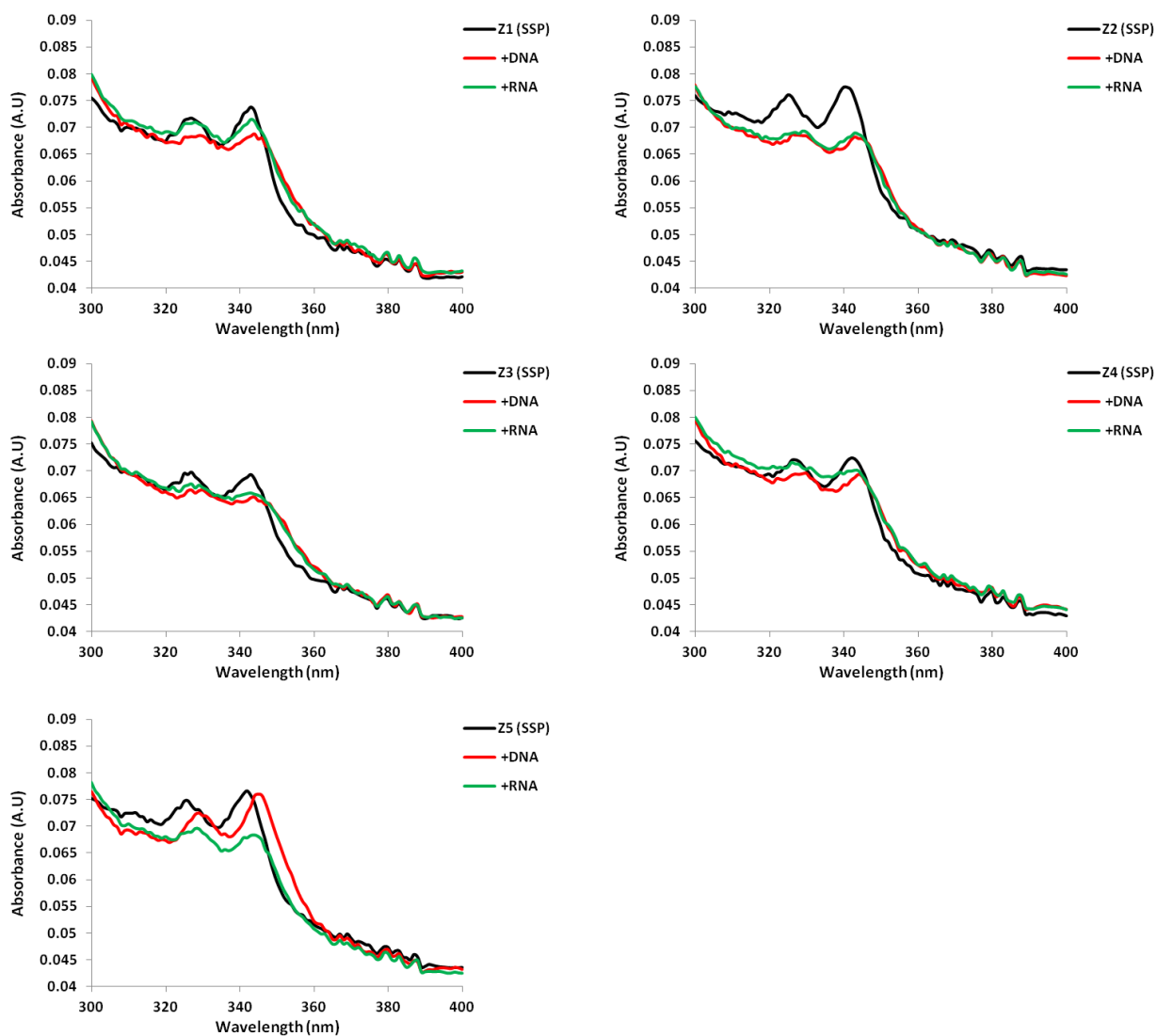


Figure S4. Absorbance spectra of single-stranded **Z1-Z5** and their corresponding duplexes with DNA/RNA targets. For conditions, see Fig. S2.

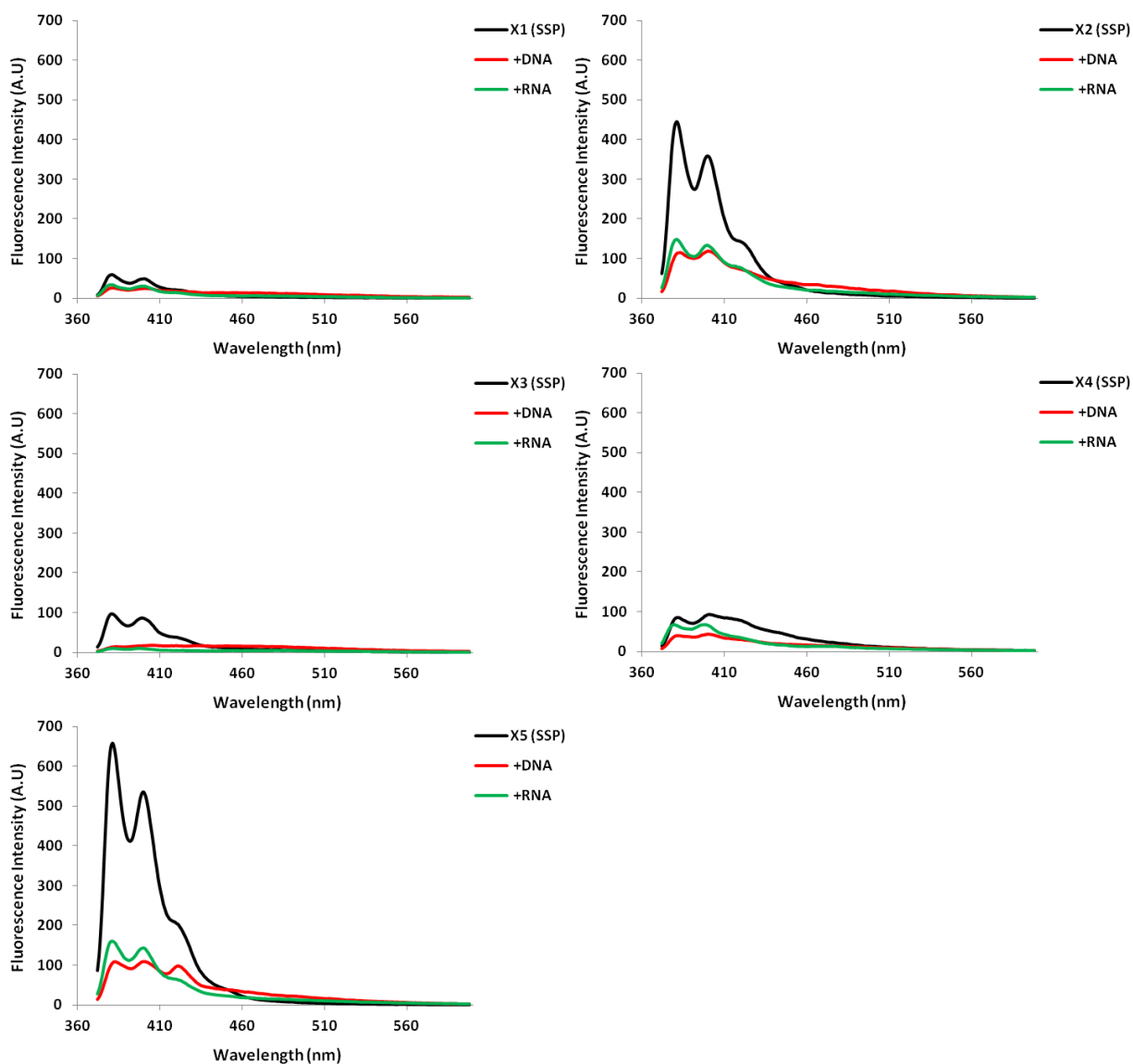


Figure S5. Steady-state fluorescence emission spectra of **X**-modified ONs and their corresponding duplexes with DNA/RNA targets. Spectra were recorded at $T = 5\text{ }^{\circ}\text{C}$ using $\lambda_{\text{ex}} = 350\text{ nm}$ and each strand at $1.0\text{ }\mu\text{M}$ concentration in T_{m} buffer. Some, but not all, of these spectra have been previously published in reference S1. Spectra are included to facilitate direct comparison.

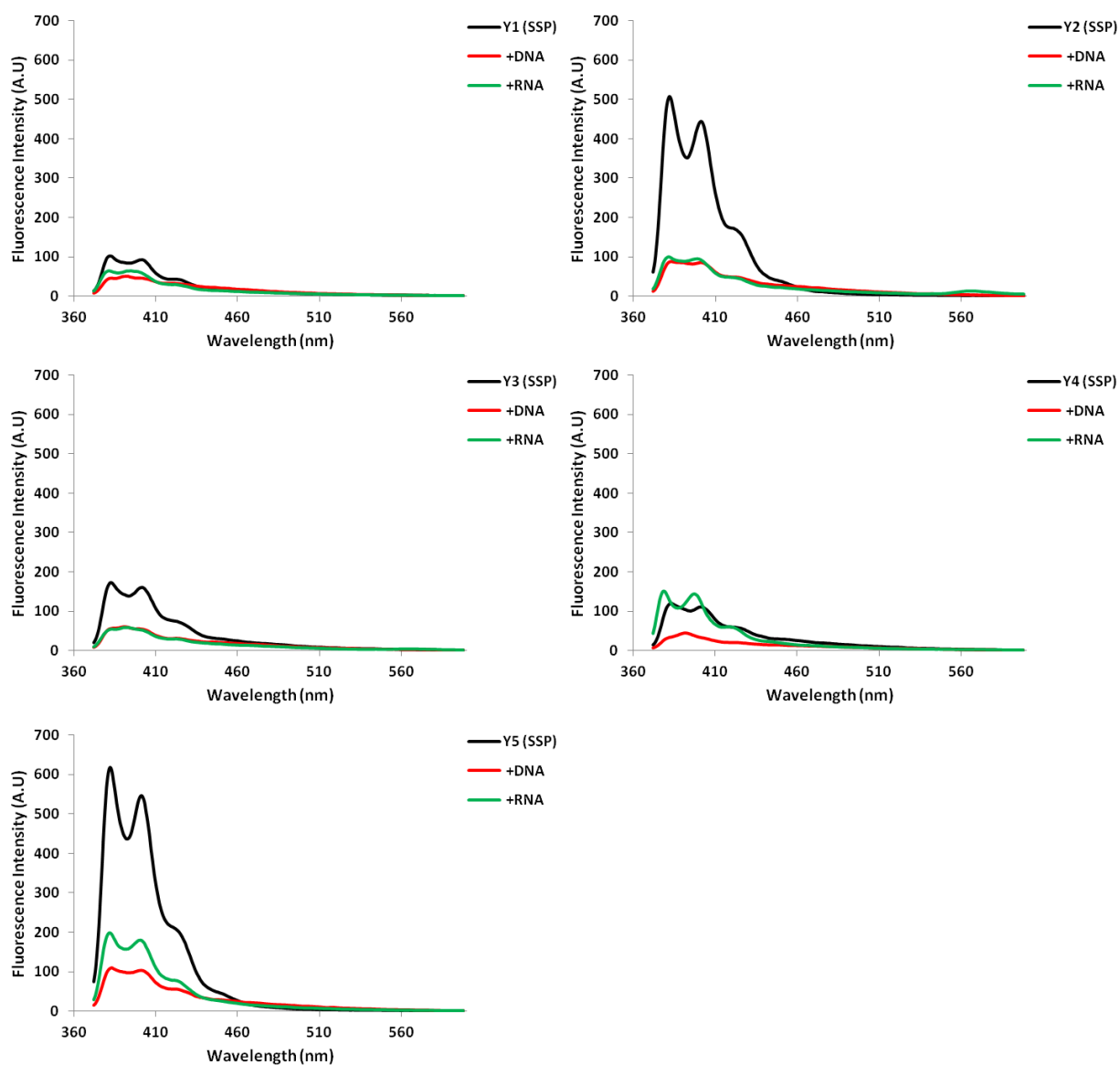


Figure S6. Steady-state fluorescence emission spectra of **Y**-modified ONs and their corresponding duplexes with DNA/RNA targets. $\lambda_{\text{ex}} = 345$ nm. For other conditions, see Fig. S5.

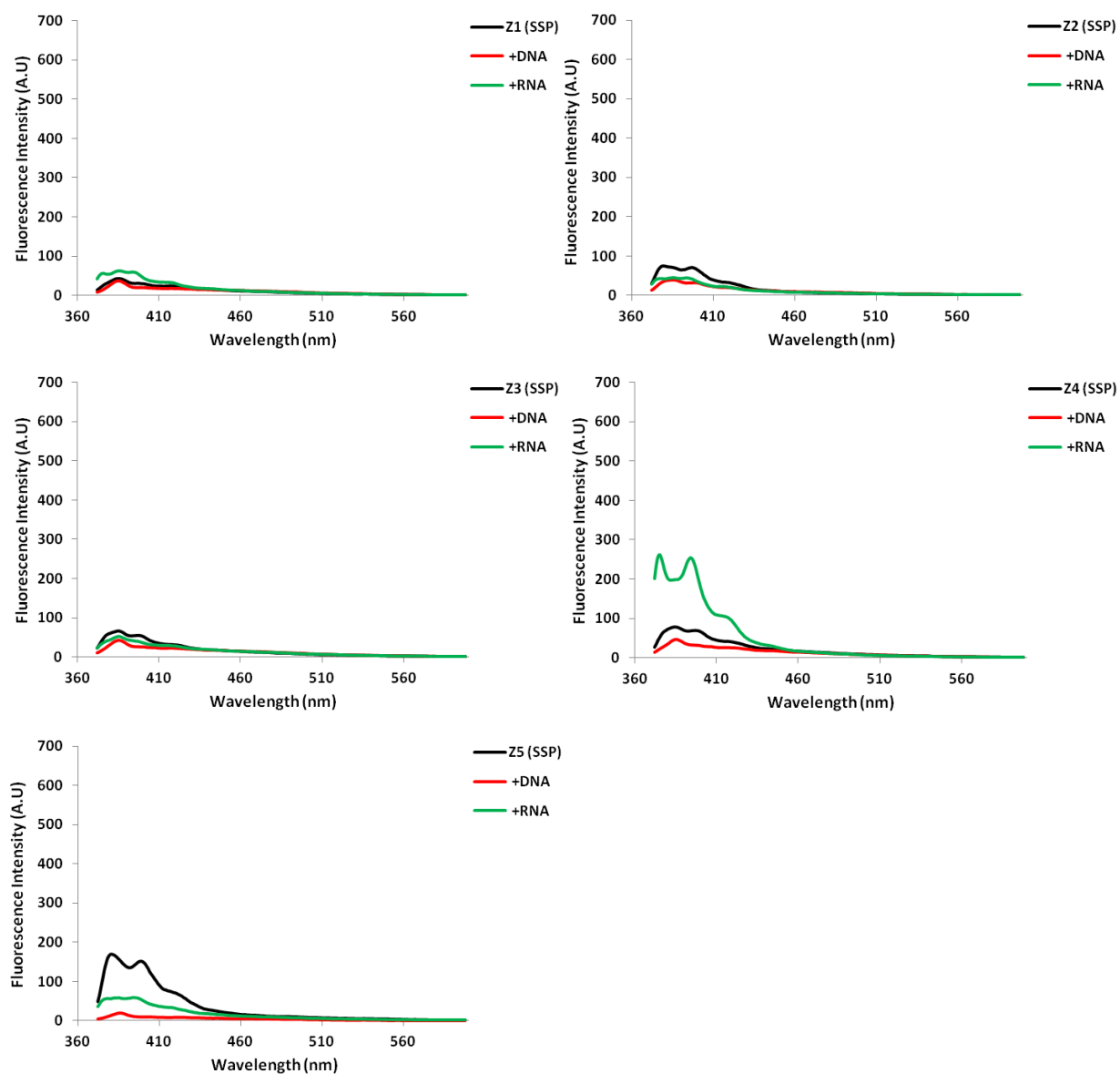


Figure S7. Steady-state fluorescence emission spectra of **Z**-modified ONs and their corresponding duplexes with DNA/RNA targets. $\lambda_{\text{ex}} = 340$ nm. For other conditions, see Fig. S5.

Table S5. Change in enthalpy upon duplex formation (ΔH) and change in enthalpy upon probe recognition of iso-sequential dsDNA target **D1:D4** (ΔH_{rec}).^a

ON	ZP	Sequence	$\Delta H^{293} [\Delta \Delta H^{293}]$ (kJ/mol)			ΔH_{rec} (kJ/mol)
			upper ON vs cDNA	lower ON vs cDNA	probe duplex	
X1 X5	+4	5'-G <u>X</u> G ATA TGC 3'-CAC T <u>A</u> X ACG	-281±4 [+25]	-299±2 [+7]	-279±1 [+27]	+5
X1 X4	+2	5'-G <u>X</u> G ATA TGC 3'-CAC <u>X</u> AT ACG	-281±4 [+25]	-300±7 [+6]	-248±8 [+58]	-27
X2 X5	+1	5'-GTG A <u>X</u> A TGC 3'-CAC T <u>A</u> X ACG	-305±1 [+1]	-299±2 [+7]	-244±3 [+62]	-54
X2 X4	-1	5'-GTG A <u>X</u> A TGC 3'-CAC <u>X</u> AT ACG	-305±1 [+1]	-300±7 [+6]	-296±6 [+10]	-3
X3 X5	-1	5'-GTG ATA <u>X</u> GC 3'-CAC T <u>A</u> X ACG	-270±7 [+36]	-299±2 [+7]	-280±5 [+26]	+17
X3 X4	-3	5'-GTG ATA <u>X</u> GC 3'-CAC <u>X</u> AT ACG	-270±7 [+36]	-300±7 [+6]	-309±4 [-3]	+45
Y1 Y5	+4	5'-G <u>Y</u> G ATA TGC 3'-CAC T <u>A</u> <u>Y</u> ACG	-292±14 [+14]	-306±15 [±0]	-277±2 [+29]	-15
Y1 Y4	+2	5'-G <u>Y</u> G ATA TGC 3'-CAC <u>Y</u> AT ACG	-292±14 [+14]	-271±20 [+35]	-238±3 [+68]	-19
Y2 Y5	+1	5'-GTG A <u>Y</u> A TGC 3'-CAC T <u>A</u> <u>Y</u> ACG	-309±10 [-3]	-306±15 [±0]	-257±5 [+49]	-52
Y2 Y4	-1	5'-GTG A <u>Y</u> A TGC 3'-CAC <u>Y</u> AT ACG	-309±10 [-3]	-271±20 [+35]	-307±6 [+1]	+31
Y3 Y5	-1	5'-GTG ATA <u>Y</u> GC 3'-CAC T <u>A</u> <u>Y</u> ACG	-317±10 [-11]	-306±15 [±0]	-288±2 [+18]	-29
Y3 Y4	-3	5'-GTG ATA <u>Y</u> GC 3'-CAC <u>Y</u> AT ACG	-317±10 [-11]	-271±20 [+35]	-276±4 [+30]	-6
Z1 Z5	+4	5'-G <u>Z</u> G ATA TGC 3'-CAC T <u>A</u> <u>Z</u> ACG	-309±8 [-3]	-253±26 [+53]	-252±4 [+54]	-4
Z1 Z4	+2	5'-G <u>Z</u> G ATA TGC 3'-CAC <u>Z</u> AT ACG	-309±8 [-3]	-332±7 [-26]	-251±5 [+55]	-84
Z2 Z5	+1	5'-GTG A <u>Z</u> A TGC 3'-CAC T <u>A</u> <u>Z</u> ACG	-260±8 [+46]	-253±26 [+53]	N/A	-
Z2 Z4	-1	5'-GTG A <u>Z</u> A TGC 3'-CAC <u>Z</u> AT ACG	-260±8 [+46]	-332±7 [-26]	-283±9 [+23]	-3
Z3 Z5	-1	5'-GTG ATA <u>Z</u> GC 3'-CAC T <u>A</u> <u>Z</u> ACG	-280±4 [+26]	-253±26 [+53]	-251±2 [+55]	+24
Z3 Z4	-3	5'-GTG ATA <u>Z</u> GC 3'-CAC <u>Z</u> AT ACG	-280±4 [+26]	-332±7 [-26]	-247±1 [+59]	-59

^a $\Delta \Delta H$ is measured relative to ΔH for **D1:D4** = -306 kJ/mol. $\Delta H_{\text{rec}} = \Delta H$ (upper strand vs cDNA) + ΔH (lower strand vs cDNA) - ΔH (probe duplex) - ΔH (dsDNA target). "±" denotes standard deviation. N/A = the lack of a clear lower base line prevented determination of this value.

Table S6. Change in entropy at 293K upon duplex formation ($-T^{293}\Delta S$) and change in entropy upon probe recognition of iso-sequential dsDNA target **D1:D4** ($-T^{293}\Delta S_{rec}$).^a

Duplex	Zipper	Sequence	$-T^{293}\Delta S$ [$\Delta(T^{293}\Delta S)$] (kJ/mol)			$-T^{293}\Delta S_{rec}$ (kJ/mol)
			upper strand vs cDNA	lower strand vs cDNA	probe duplex	
X1 X5	+4	5'-G <u>X</u> G ATA TGC 3'-CAC T <u>A</u> X ACG	235±3 [-30]	247±6 [-18]	224±1 [-41]	-7
X1 X4	+2	5'-G <u>X</u> G ATA TGC 3'-CAC <u>X</u> AT ACG	235±3 [-30]	254±6 [-11]	203±8 [-62]	+21
X2 X5	+1	5'-GTG A <u>X</u> A TGC 3'-CAC T <u>A</u> X ACG	251±1 [-14]	247±6 [-18]	204±3 [-61]	+29
X2 X4	-1	5'-GTG A <u>X</u> A TGC 3'-CAC <u>X</u> AT ACG	251±1 [-14]	254±6 [-11]	245±5 [-20]	-5
X3 X5	-1	5'-GTG ATA <u>X</u> GC 3'-CAC T <u>A</u> X ACG	220±7 [-45]	247±6 [-18]	225±5 [-40]	-23
X3 X4	-3	5'-GTG ATA <u>X</u> GC 3'-CAC <u>X</u> AT ACG	220±7 [-45]	254±6 [-11]	256±3 [-9]	-47
Y1 Y5	+4	5'-G <u>Y</u> G ATA TGC 3'-CAC T <u>A</u> Y ACG	244±14 [-21]	251±14 [-14]	221±2 [-44]	+9
Y1 Y4	+2	5'-G <u>Y</u> G ATA TGC 3'-CAC <u>Y</u> AT ACG	244±14 [-21]	225±20 [-40]	193±3 [-72]	-3
Y2 Y5	+1	5'-GTG A <u>Y</u> A TGC 3'-CAC T <u>A</u> Y ACG	253±10 [-12]	251±14 [-14]	214±5 [-51]	+25
Y2 Y4	-1	5'-GTG A <u>Y</u> A TGC 3'-CAC <u>Y</u> AT ACG	253±10 [-12]	225±20 [-40]	252±5 [-13]	-39
Y3 Y5	-1	5'-GTG ATA <u>Y</u> GC 3'-CAC T <u>A</u> Y ACG	266±11 [+1]	251±14 [-14]	230±2 [-35]	+22
Y3 Y4	-3	5'-GTG ATA <u>Y</u> GC 3'-CAC <u>Y</u> AT ACG	266±11 [+1]	225±20 [-40]	223±4 [-42]	+3
Z1 Z5	+4	5'-G <u>Z</u> G ATA TGC 3'-CAC T <u>A</u> Z ACG	265±8 [±0]	205±24 [-60]	203±3 [-62]	+2
Z1 Z4	+2	5'-G <u>Z</u> G ATA TGC 3'-CAC <u>Z</u> AT ACG	265±8 [±0]	291±7 [+26]	210±5 [-55]	+81
Z2 Z5	+1	5'-GTG A <u>Z</u> A TGC 3'-CAC T <u>A</u> Z ACG	212±8 [-53]	205±24 [-60]	N/A	N/A
Z2 Z4	-1	5'-GTG A <u>Z</u> A TGC 3'-CAC <u>Z</u> AT ACG	212±8 [-53]	291±7 [+26]	242±11 [-23]	-4
Z3 Z5	-1	5'-GTG ATA <u>Z</u> GC 3'-CAC T <u>A</u> Z ACG	234±3 [-31]	205±24 [-60]	205±2 [-60]	-31
Z3 Z4	-3	5'-GTG ATA <u>Z</u> GC 3'-CAC <u>Z</u> AT ACG	234±3 [-31]	291±7 [+26]	205±1 [-60]	+55

^a $\Delta(T^{293}\Delta S)$ is measured relative to $-T^{293}\Delta S$ for **D1:D4** = 265 kJ/mol. $-T^{293}\Delta S_{rec} = T^{293}\Delta S$ (upper strand vs cDNA) + $T^{293}\Delta S$ (lower strand vs cDNA) - $T^{293}\Delta S$ (probe duplex) - $T^{293}\Delta S$ (dsDNA target). “±” denotes standard deviation. N/A = the lack of a clear lower base line prevented determination of this value.

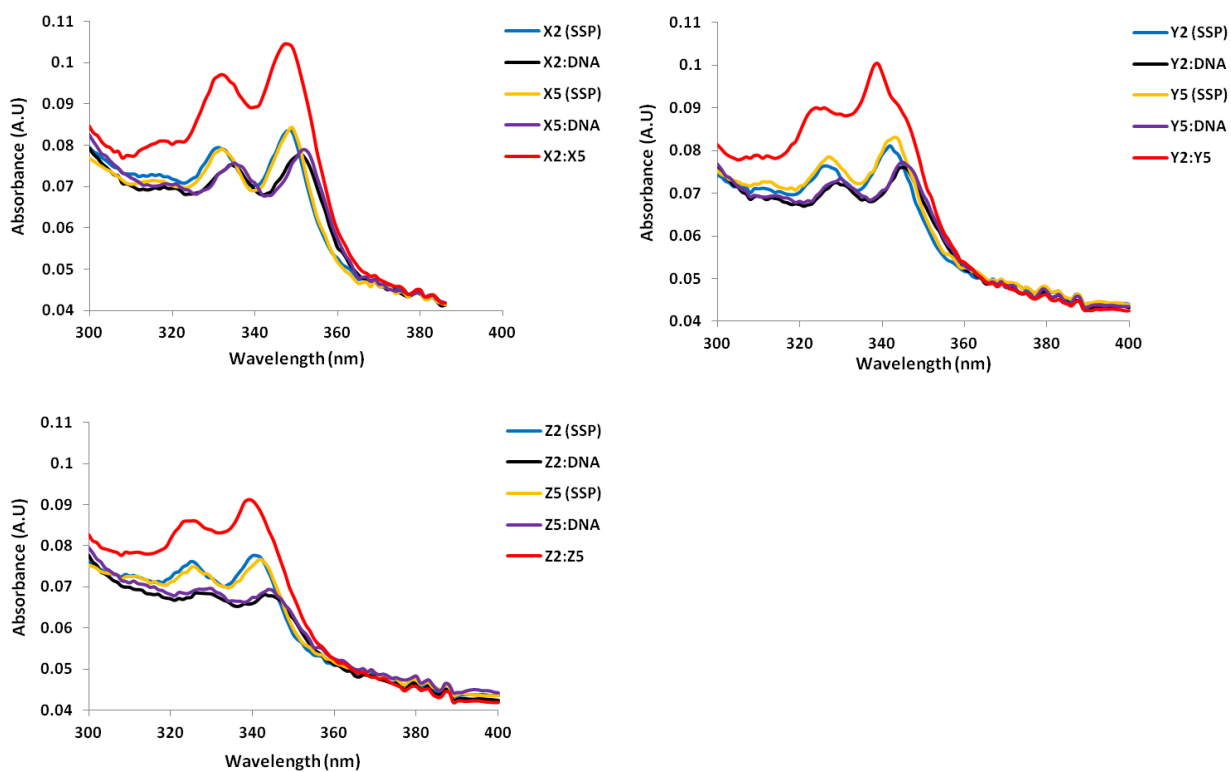


Figure S8. Absorbance spectra of representative Invaders, duplexes between individual probe strands and cDNA, and single-stranded probes (SSP). Spectra were recorded at $T = 5\text{ }^{\circ}\text{C}$ using each strand at $1.0\text{ }\mu\text{M}$ concentration in T_m buffer.

Table S7. Pseudorotational phase angles P of selected nucleotides in **B2:D4** duplexes.

P	D1:D4		X2:D4		Y2:D4		Z2:D4	
A ₄ :T ₁₅	93	112	96	116	105	110	102	106
B ₅ :A ₁₄	129	111	142	98	148	103	149	175
A ₆ :B ₁₃	102	109	124	97	155	113	171	141
T ₇ :A ₁₂	121	123	90	100	106	102	111	98

Table S8. Pseudorotational phase angles P of selected nucleotides in **B2:B5** duplexes.

P	D1:D4		X2:X5				Y2:Y5		Z2:Z5			
			Type A		Type B		Type A		Type A		Type B	
A ₄ :T ₁₅	93	112	70	103	100	104	101	106	98	105	100	106
B ₅ :A ₁₄	129	111	161 ^a	122	149	168	148	104	147	107	150	180
A ₆ :B ₁₃	102	109	123	160	176	148	104	147	107	146	173	148 ^a
T ₇ :A ₁₂	121	123	104	71	106	116	109	117	105	110	106	111

^a Not intercalated.



Figure S9. Structural base pair parameters for duplexes **D1:D4** and **B2:D4**.



Figure S10. Structural dinucleotide step parameters for duplexes **D1:D4** and **B2:D4**.

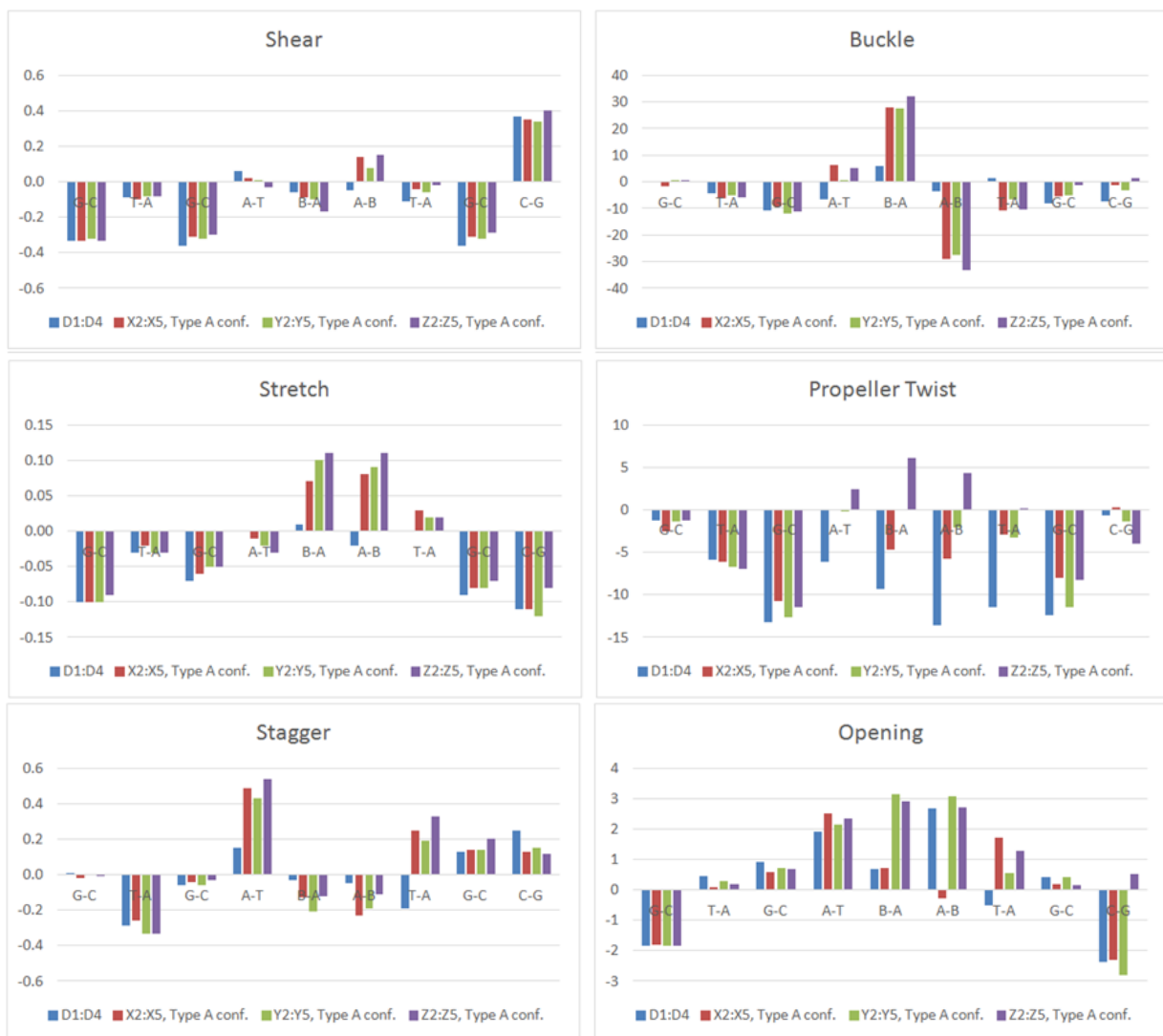


Figure S11. Structural base pair parameters for duplexes **D1:D4** and **B2:B5** (Type A).



Figure S12. Structural dinucleotide step parameters for duplexes **D1:D4** and **B2:B5** (Type A).

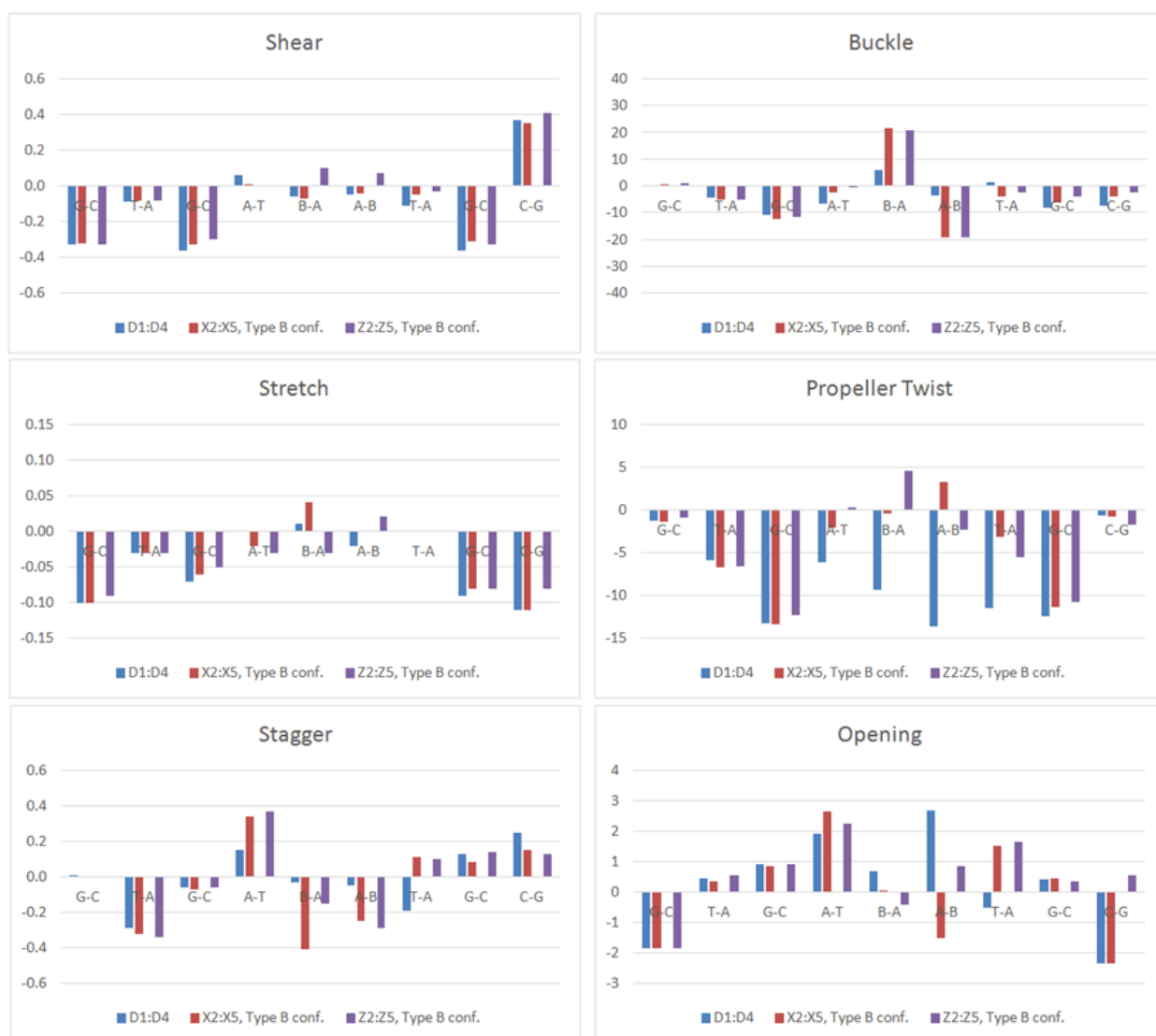


Figure S13. Structural base pair parameters for duplexes **D1:D4** and **B2:B5** (Type B).

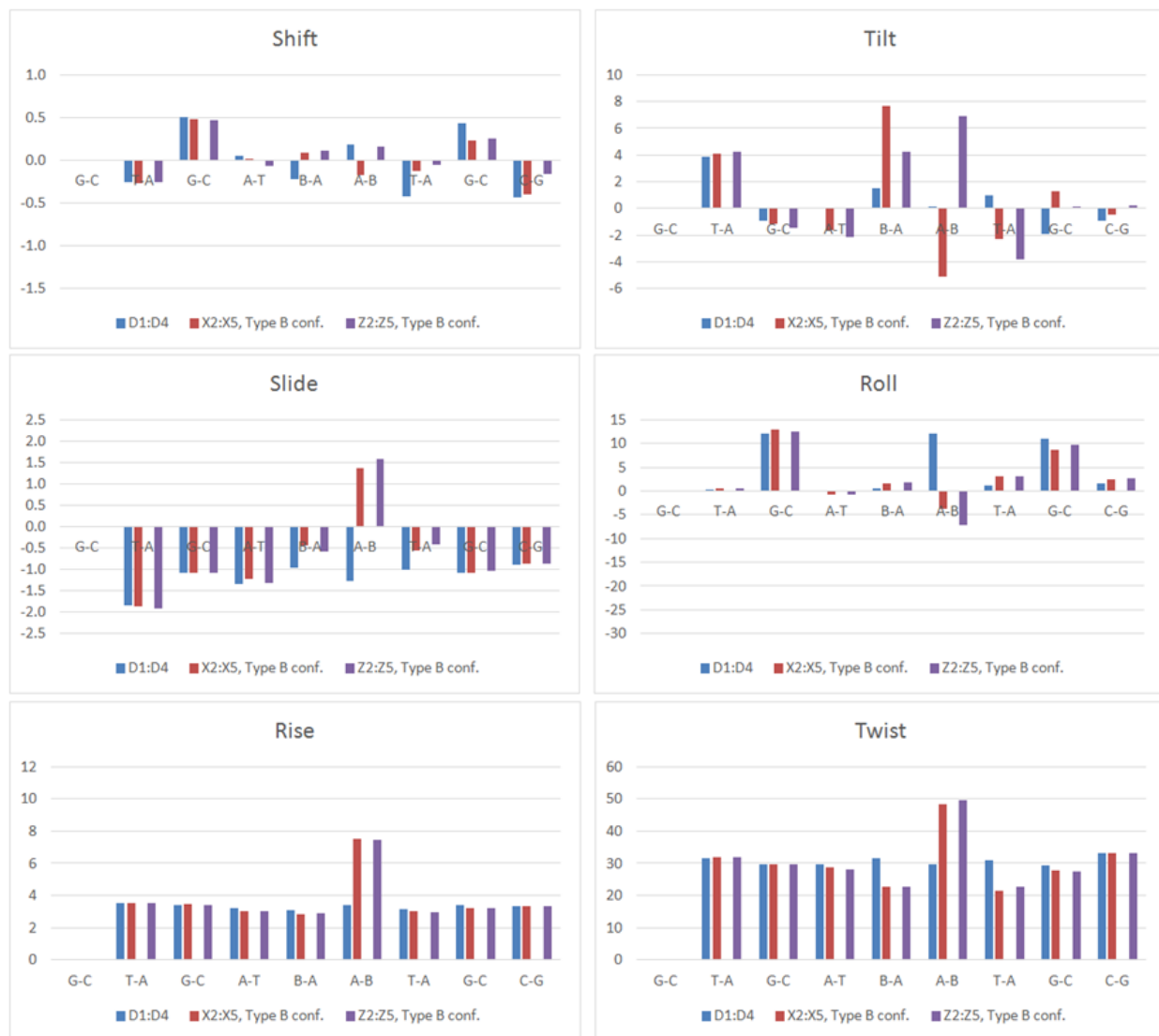


Figure S14. Structural dinucleotide step parameters for duplexes **D1:D4** and **B2:B5** (Type B).

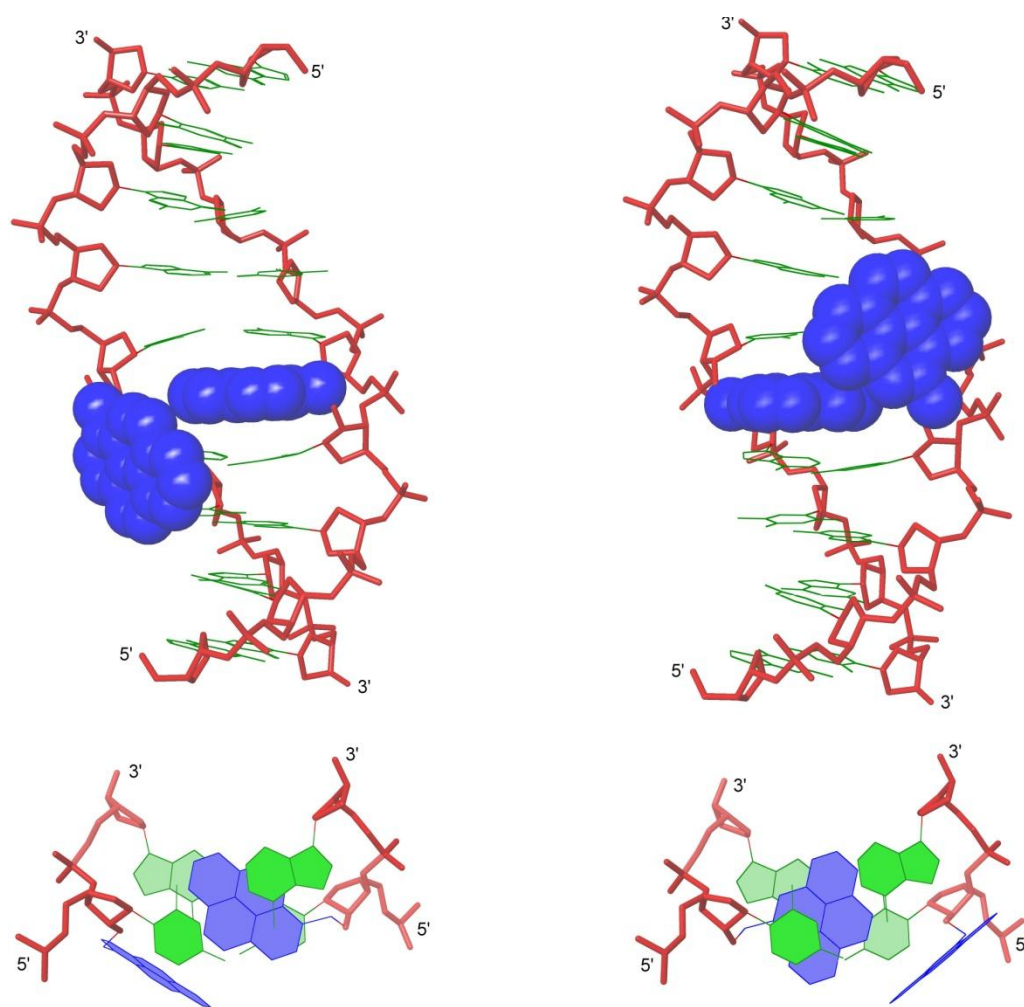
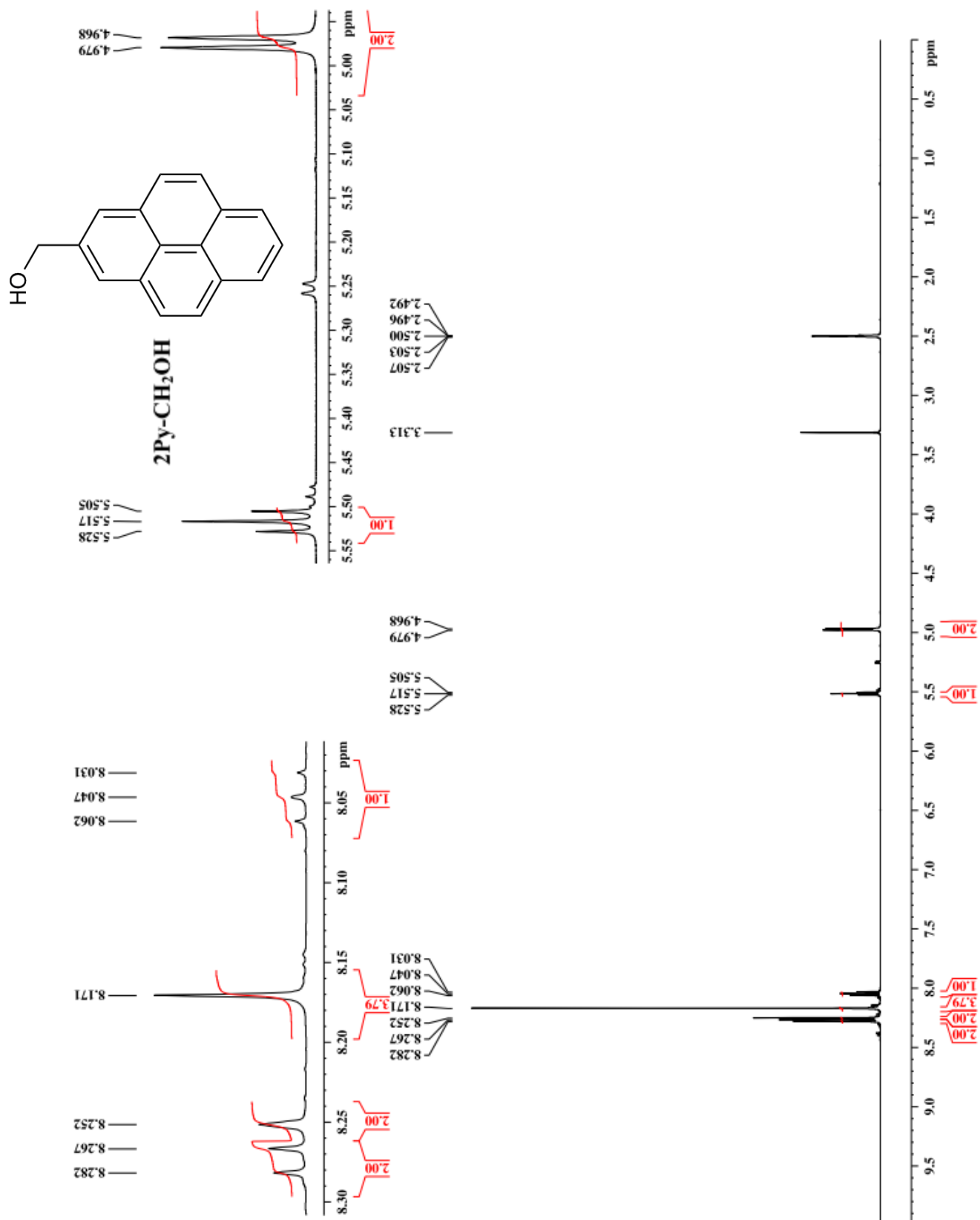
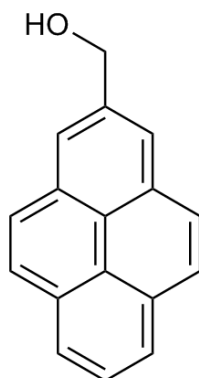


Figure S15. Lowest energy structures of **X2:X5** (left) and **Z2:Z5** (right) in Type B conformation (**Y2:Y5** maintained a Type A conformation throughout the simulation protocol). Top: side view of duplex; bottom: top view of the central duplex region. Color code: sugar phosphate backbone (red); pyrene moieties (blue) and nucleobases (green). Hydrogen atoms, sodium ions and bond orders are omitted for clarity.

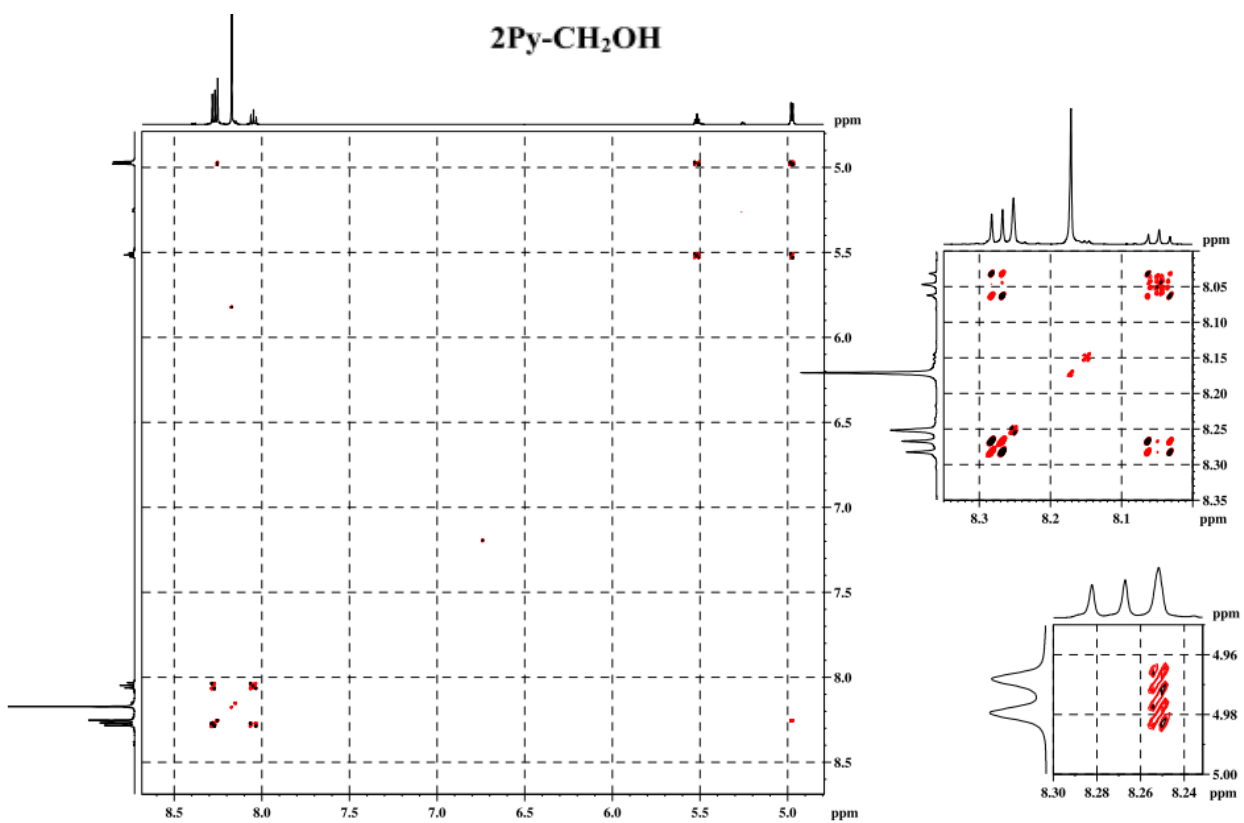
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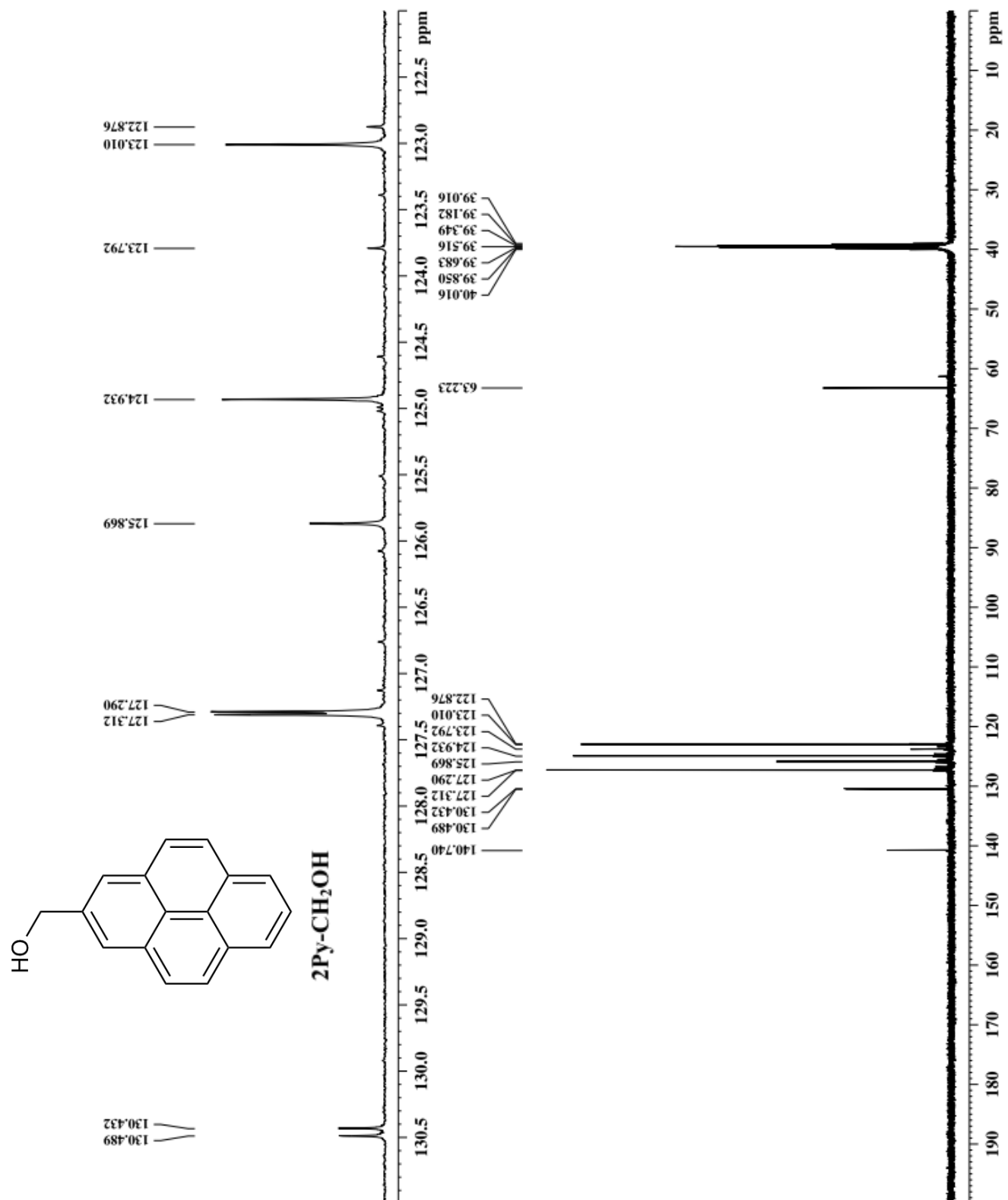
(S1) S. Karmakar, B. A. Anderson, R. L. Rathje, S. Andersen, T. Jensen, P. Nielsen and P. J. Hrdlicka, *J. Org. Chem.*, 2011, **76**, 7119-7131.

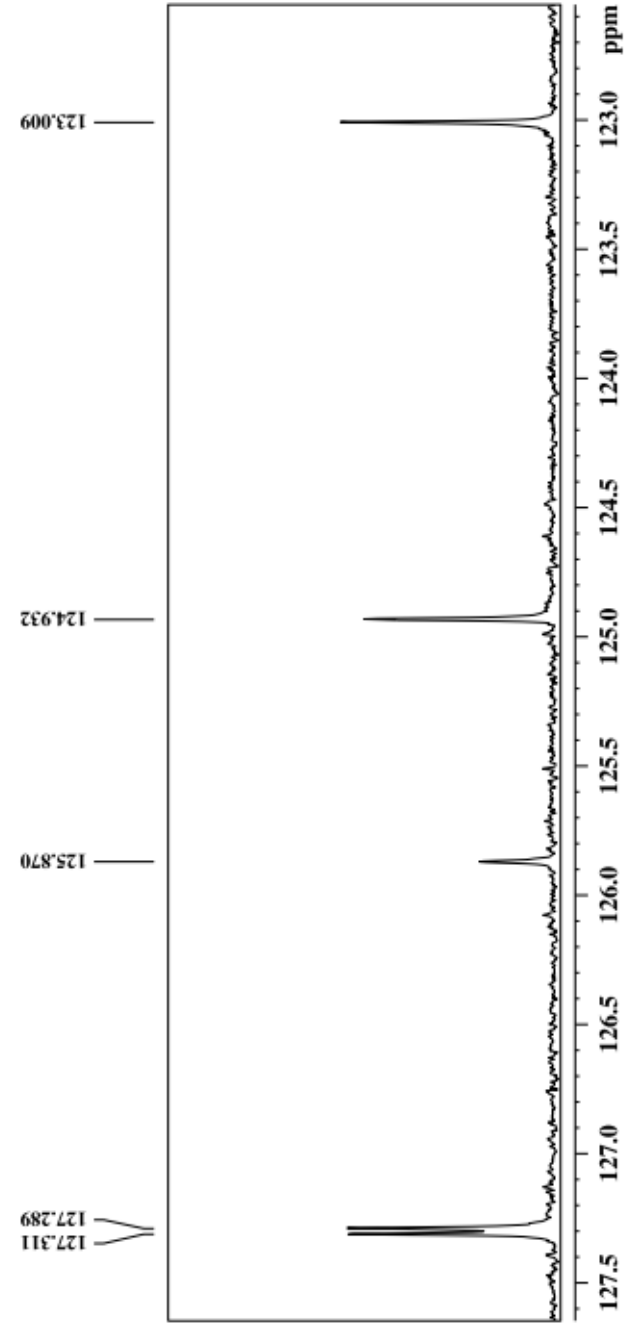
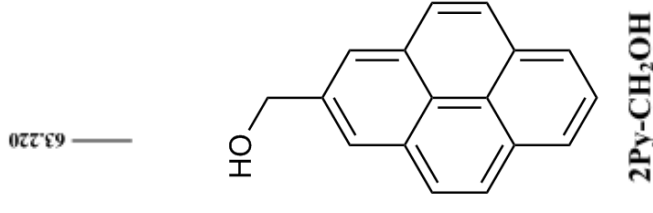




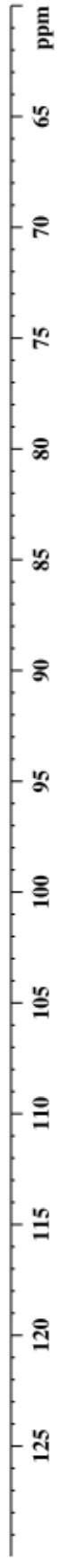
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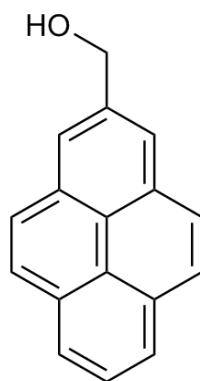




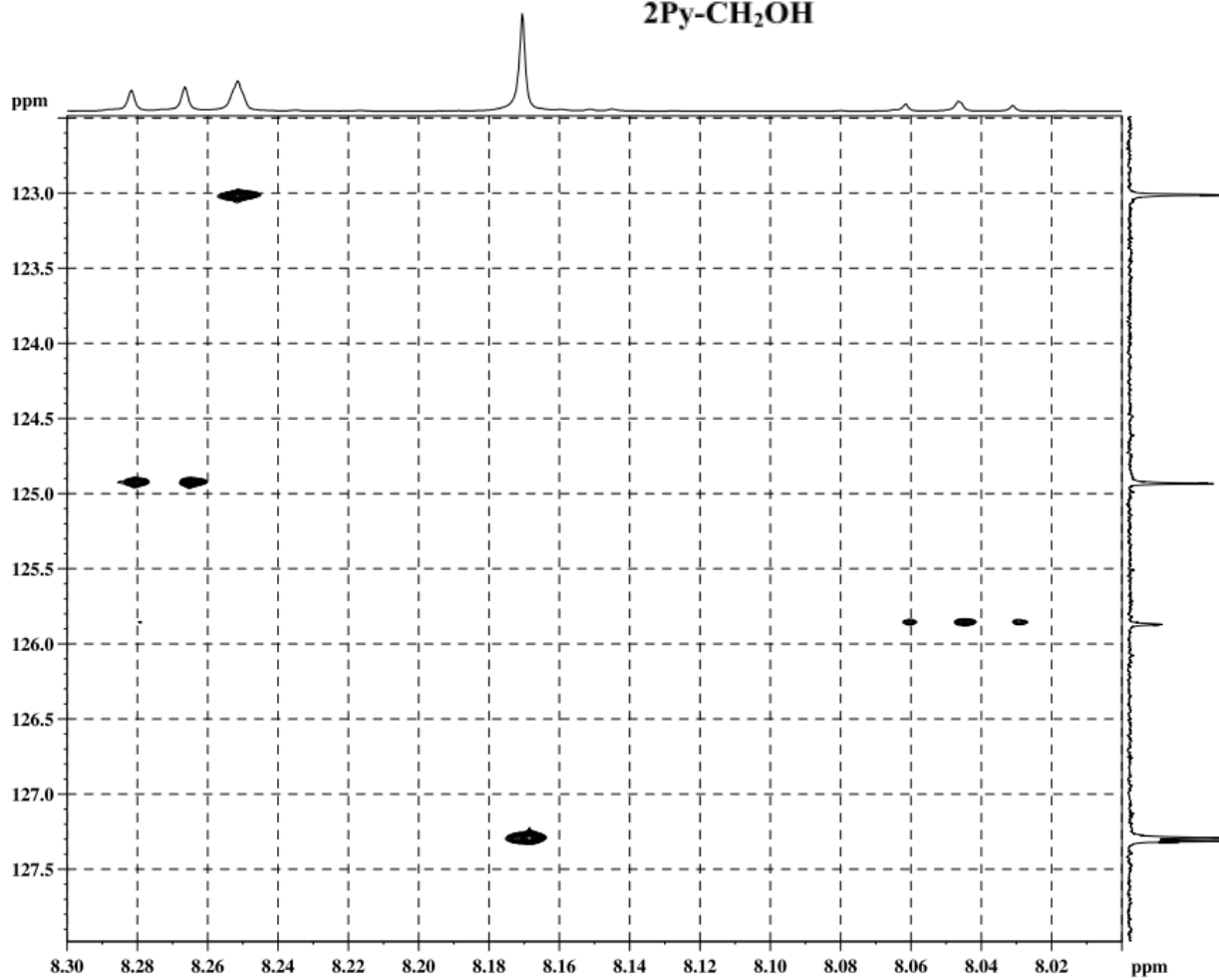


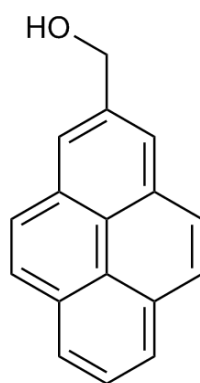
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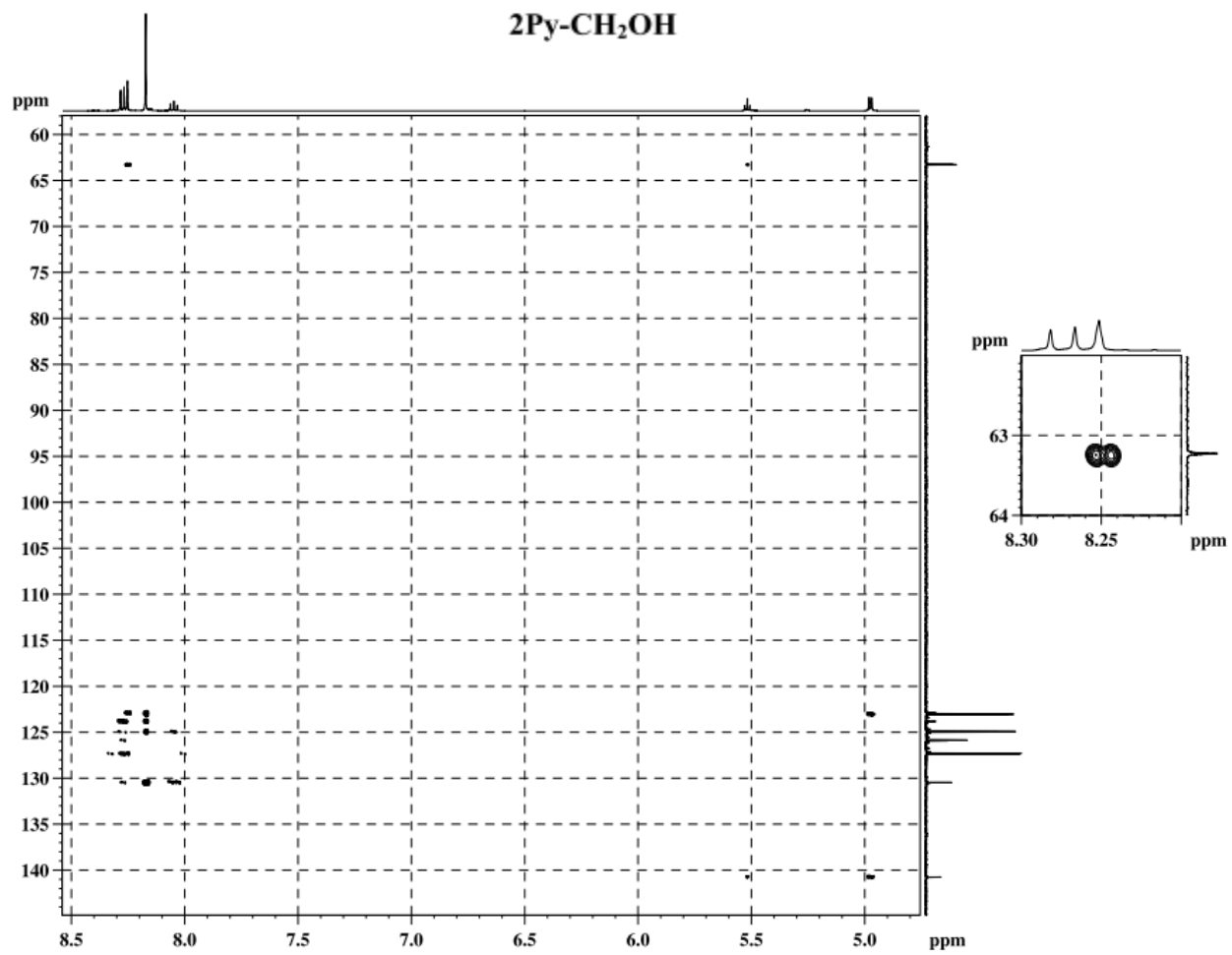


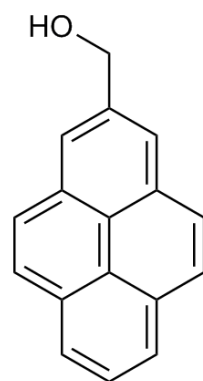
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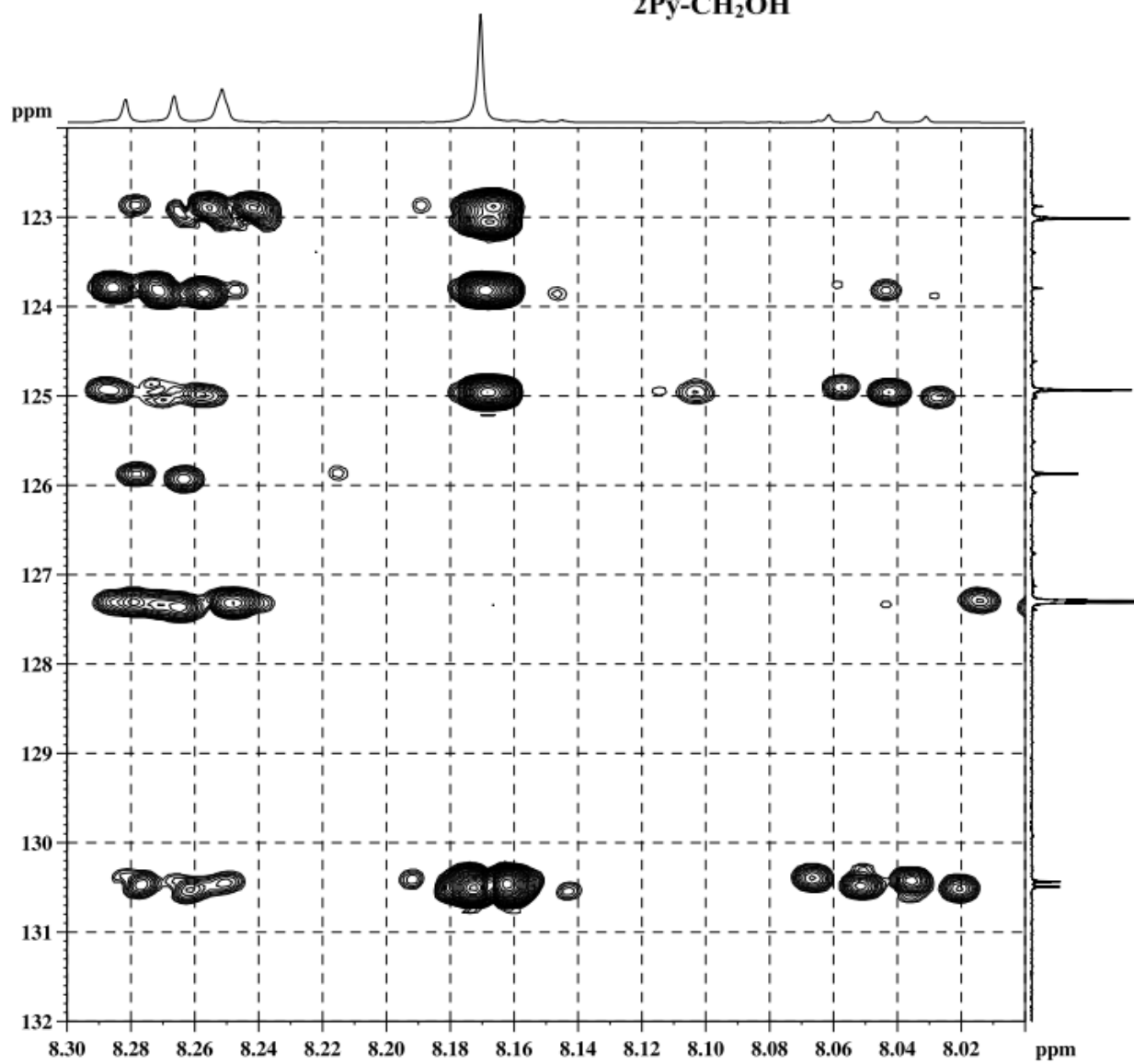


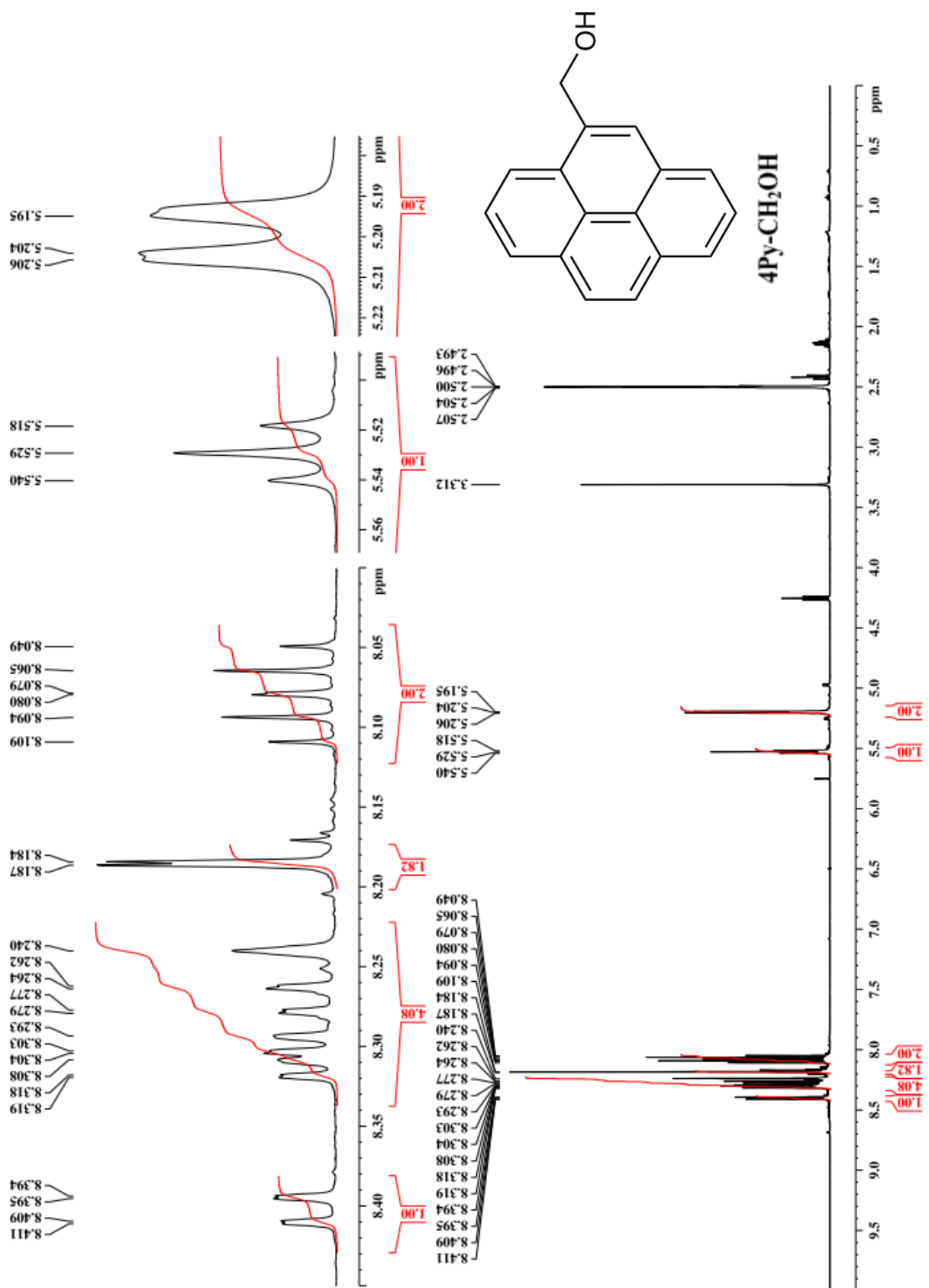
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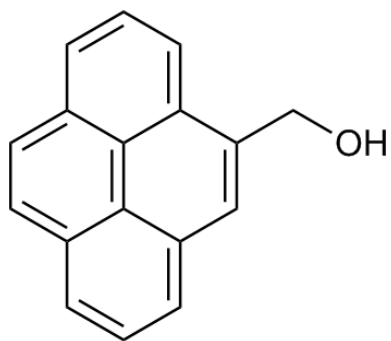




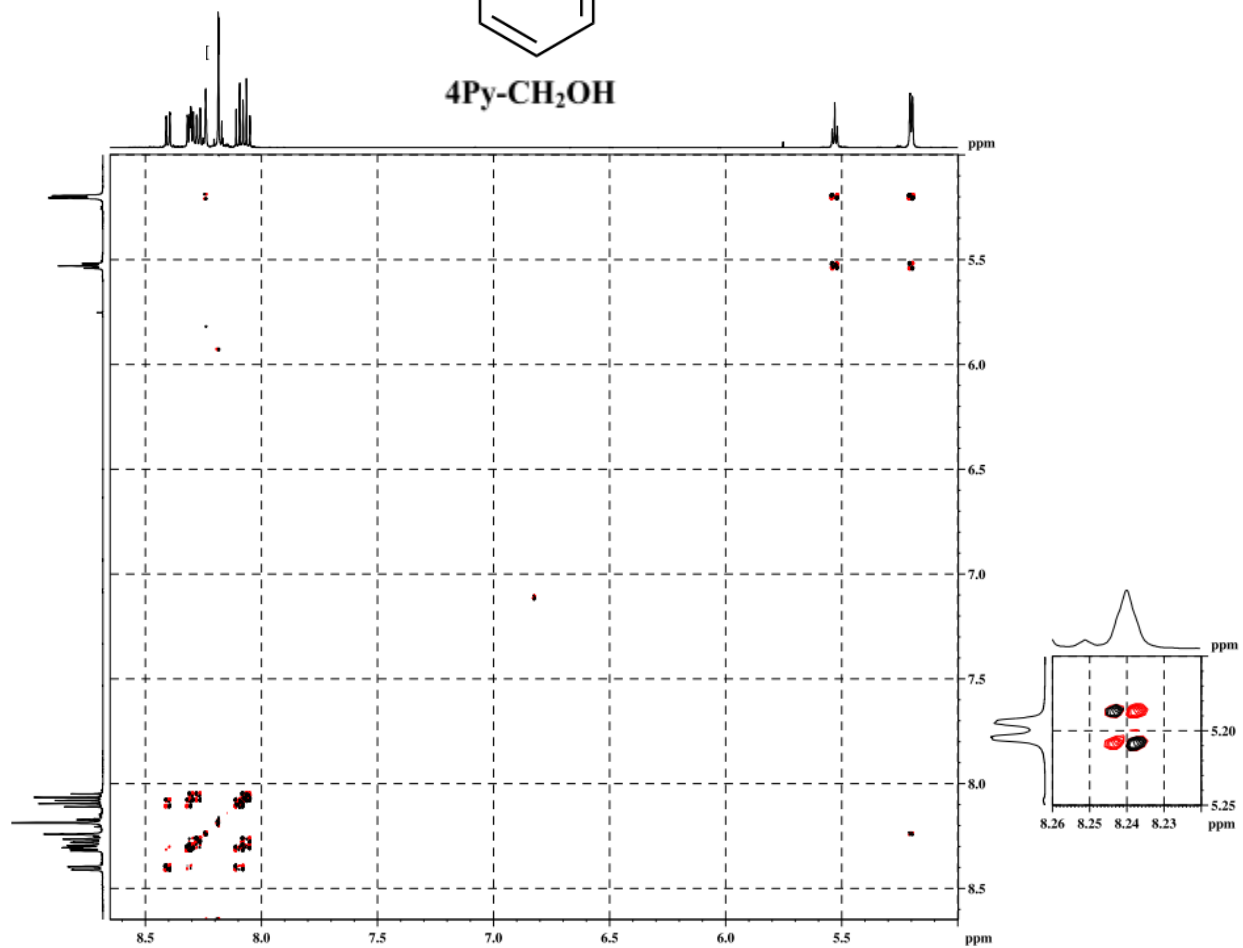
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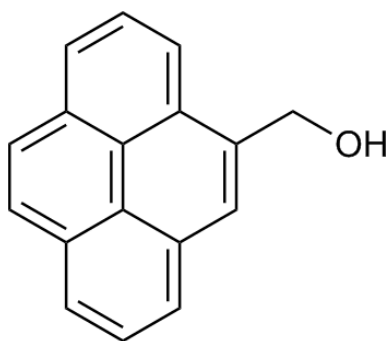




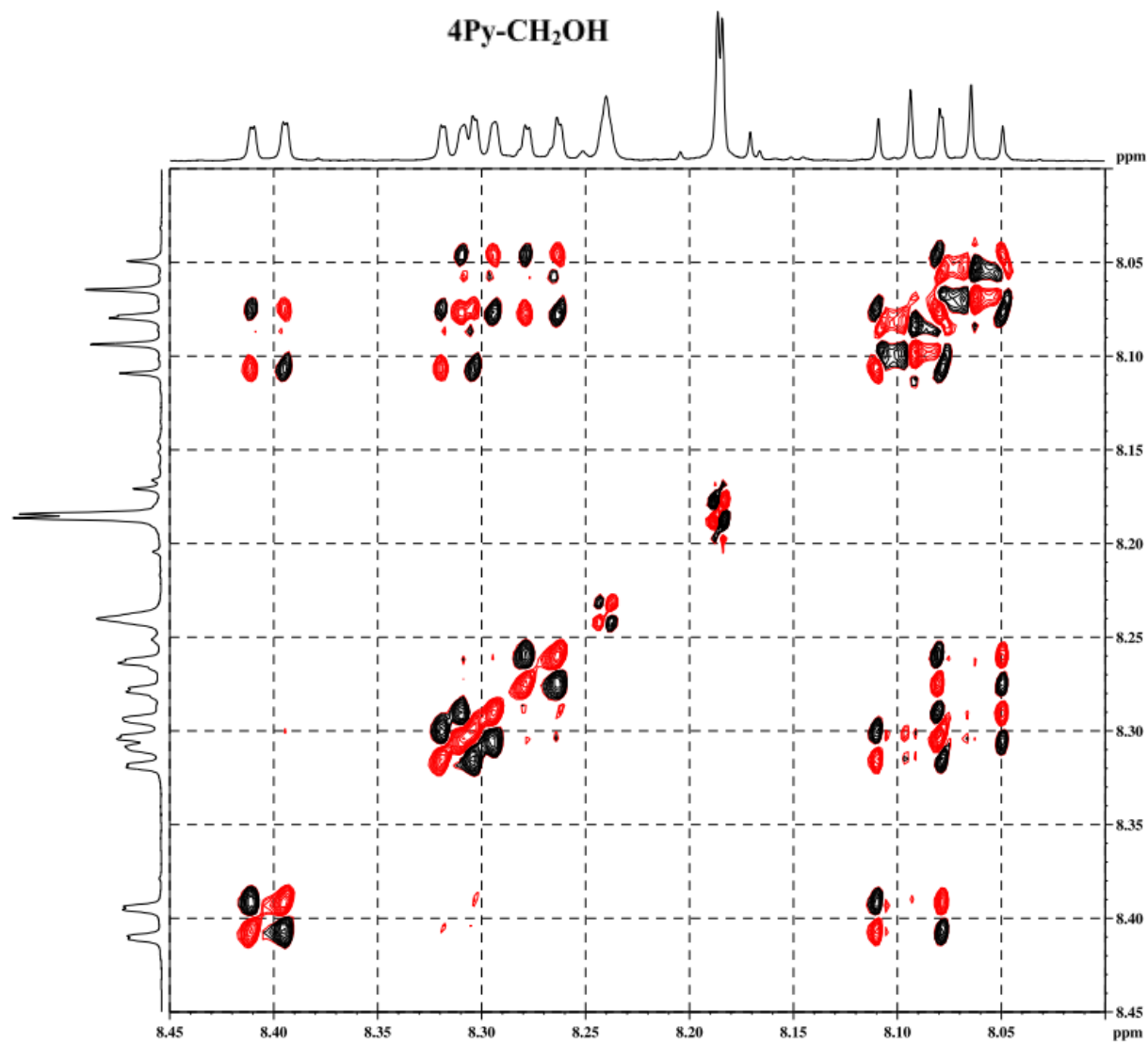


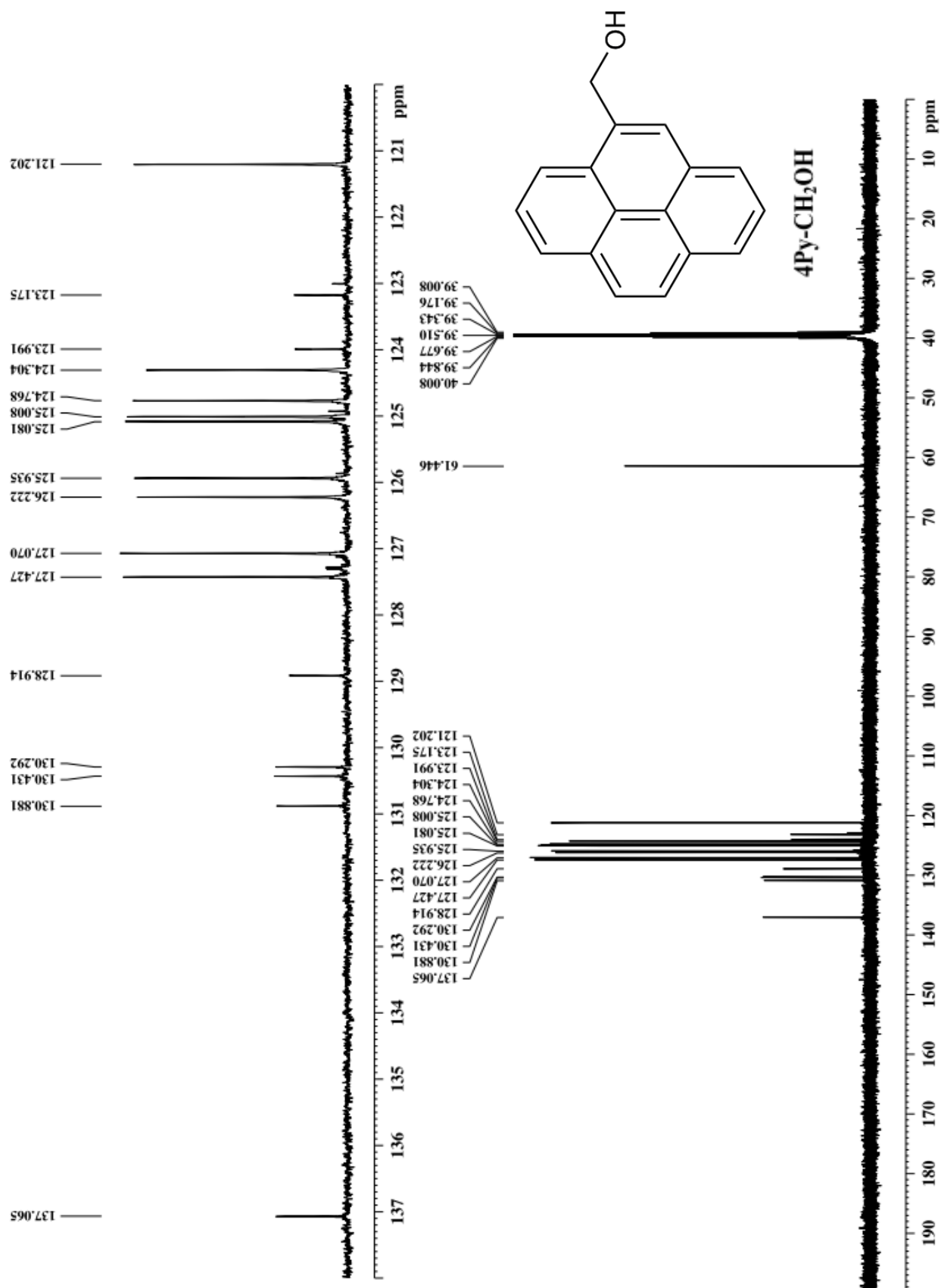
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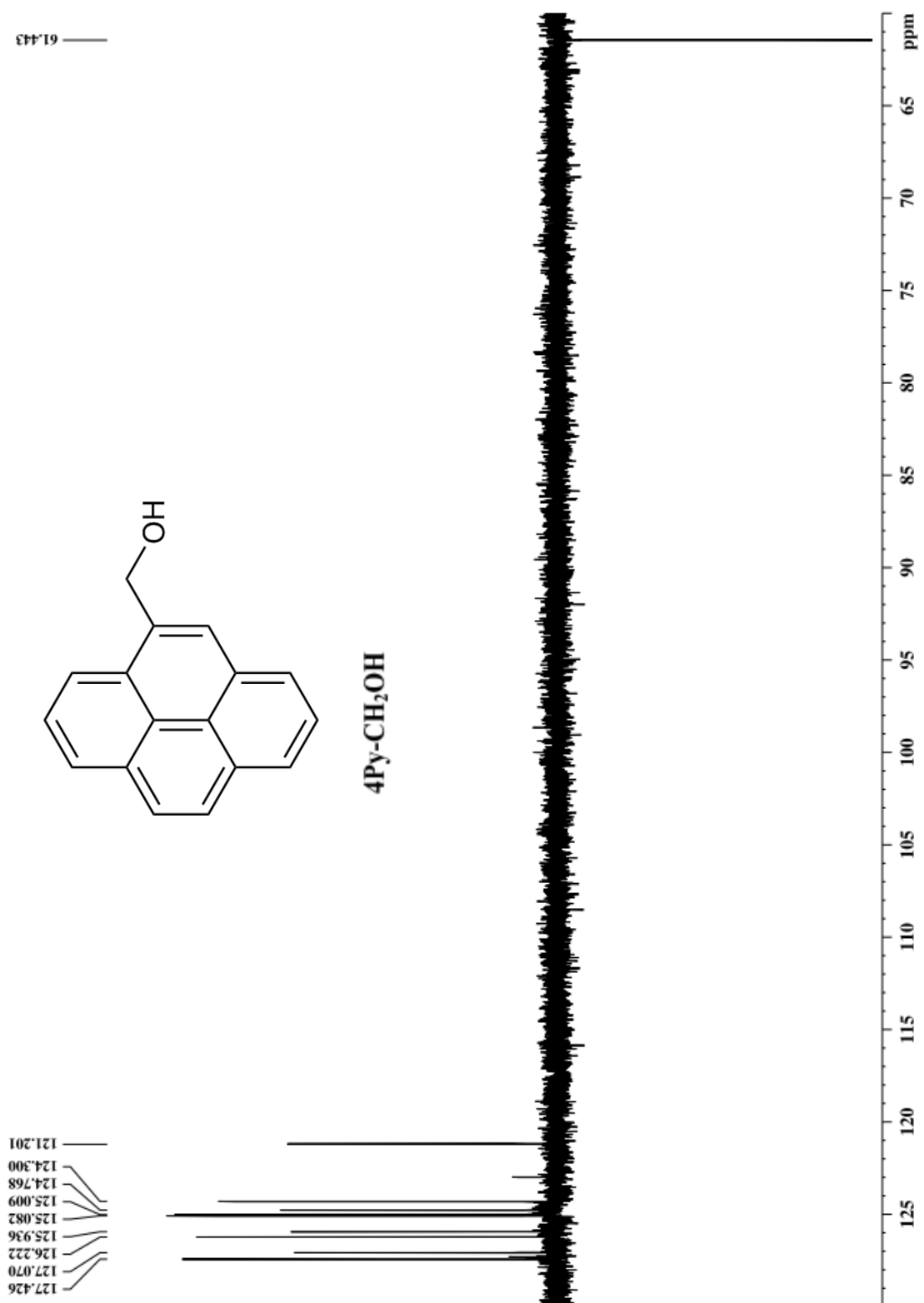


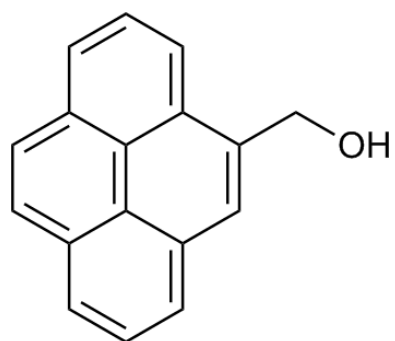


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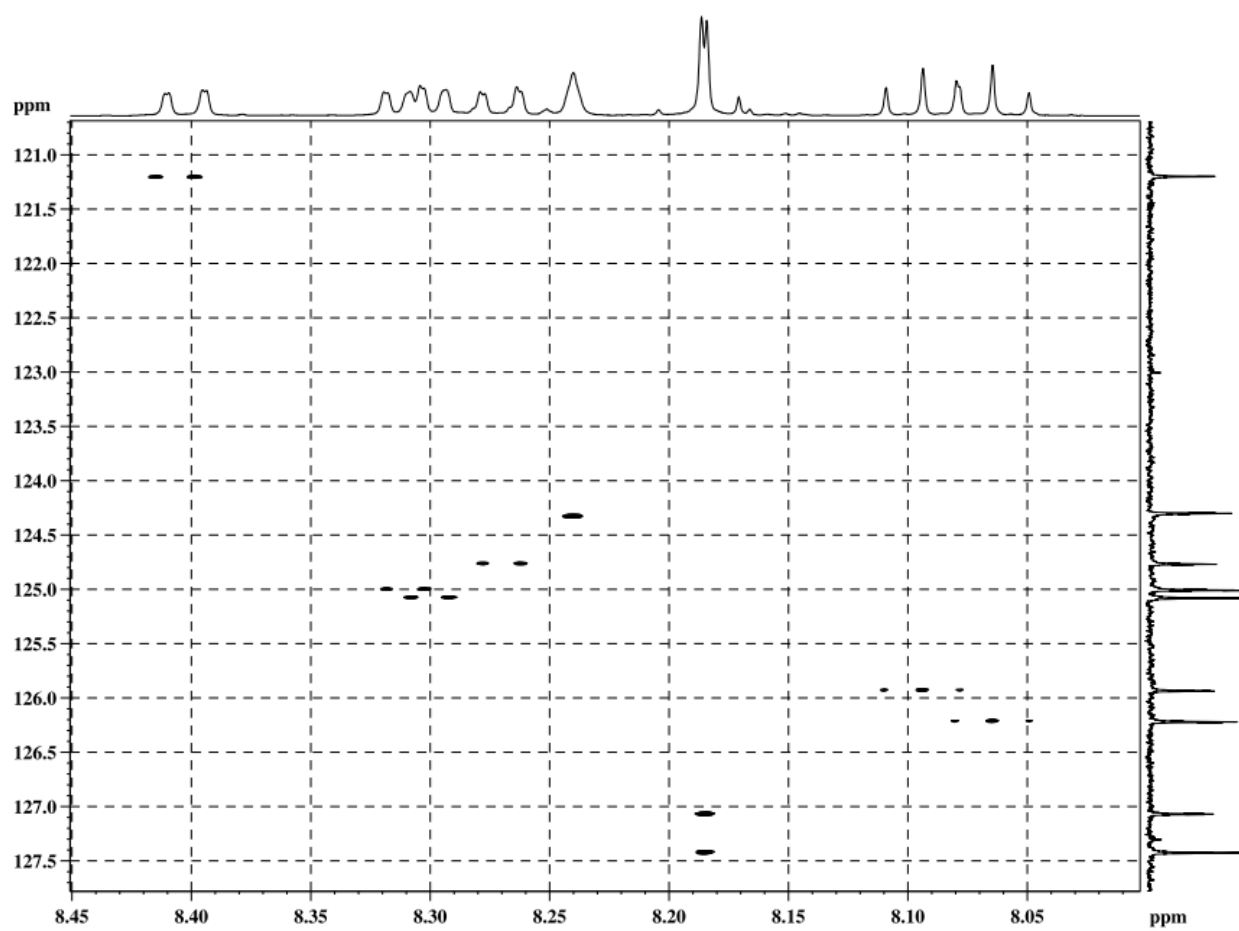


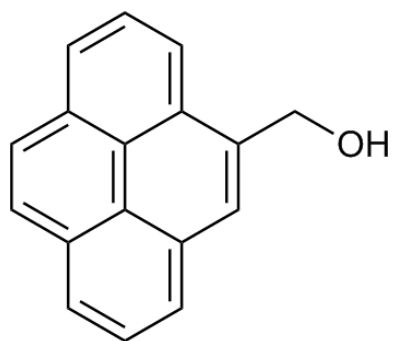




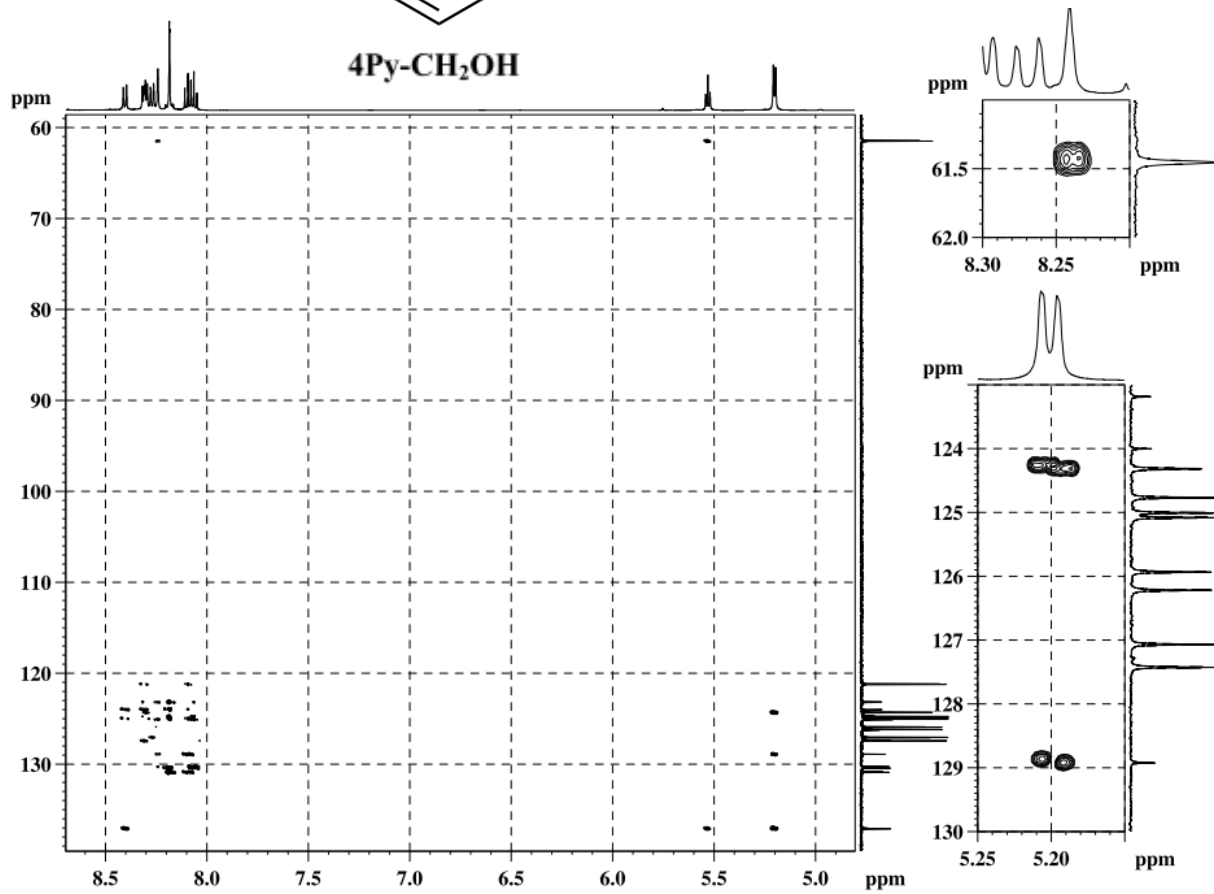


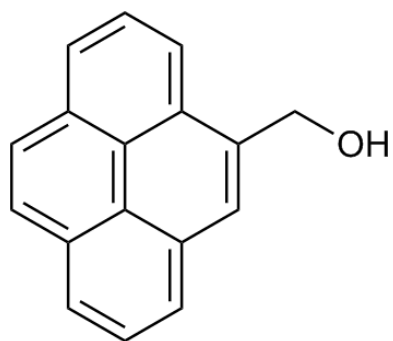
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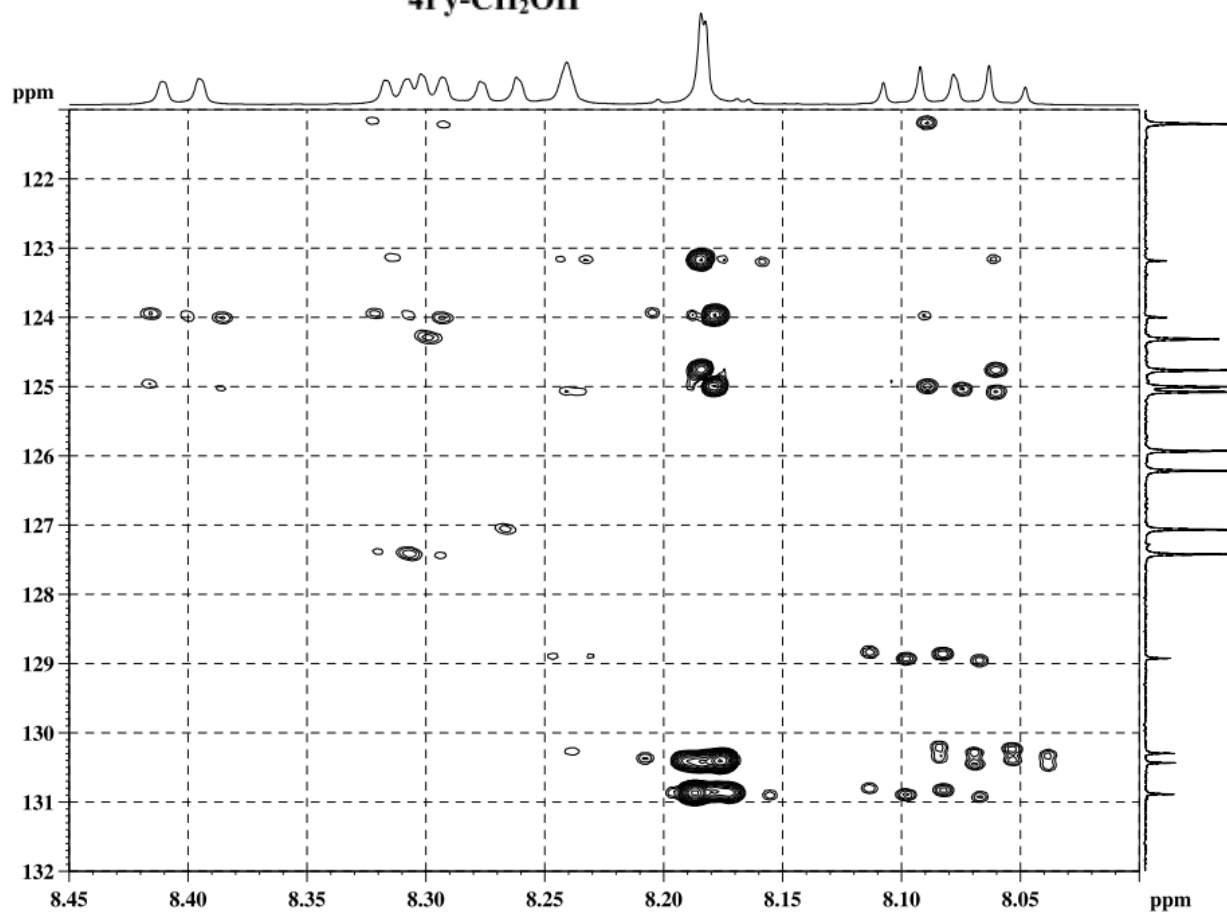


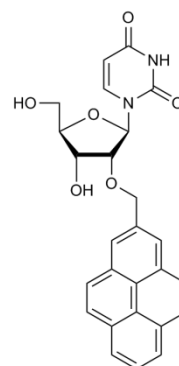
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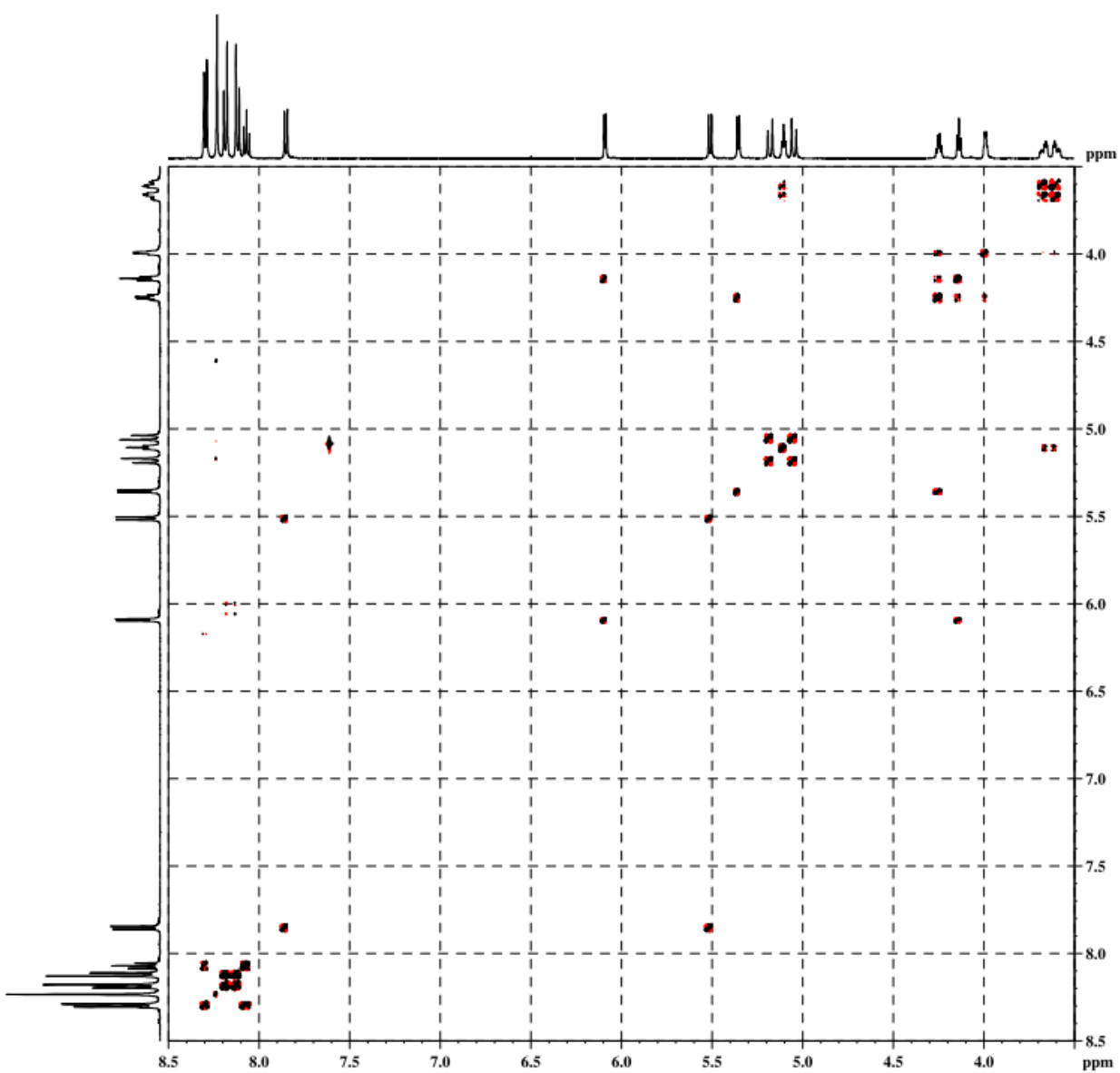


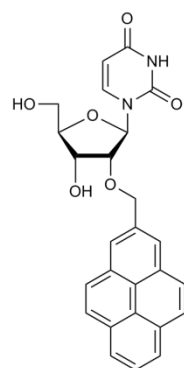
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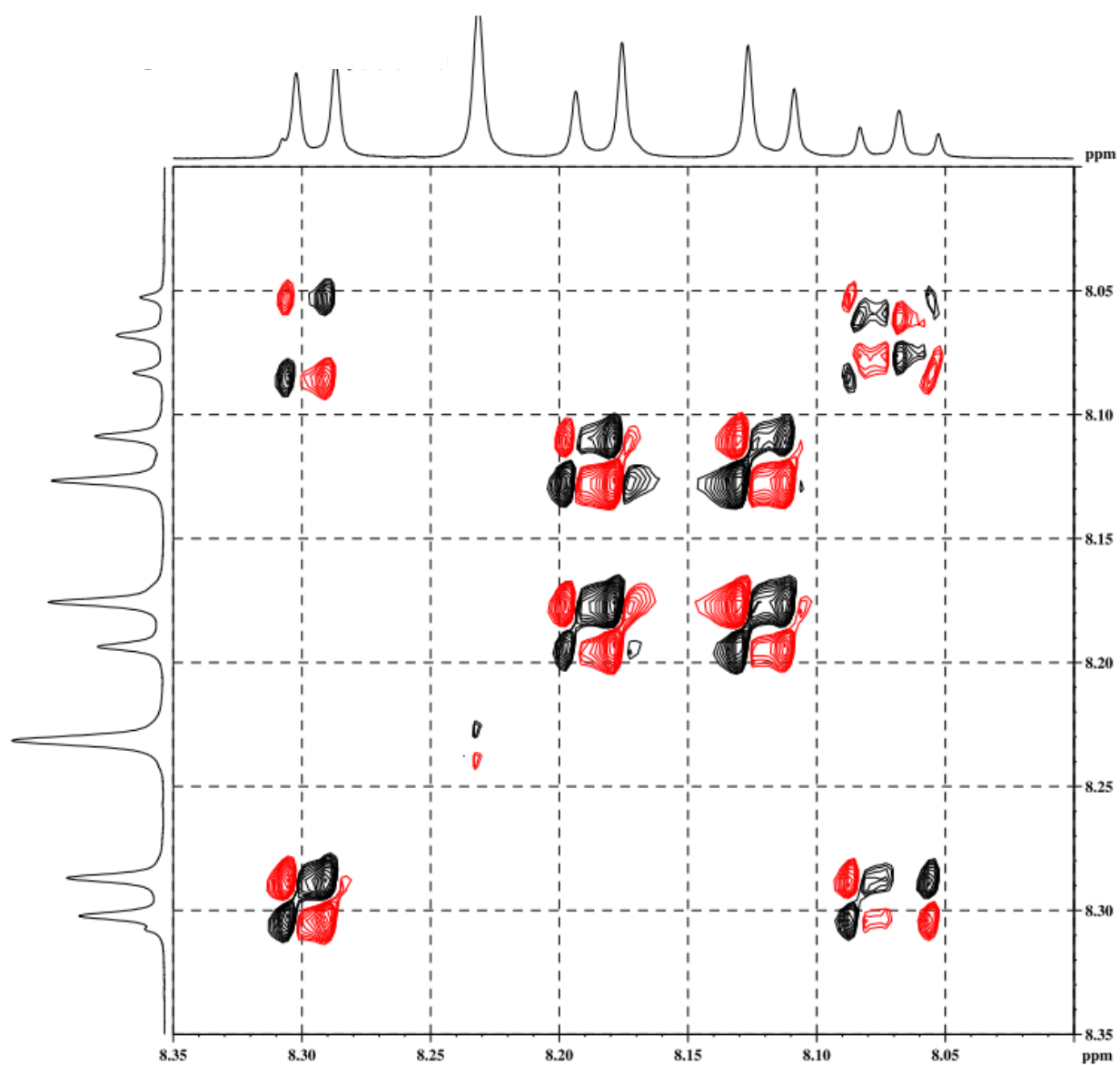


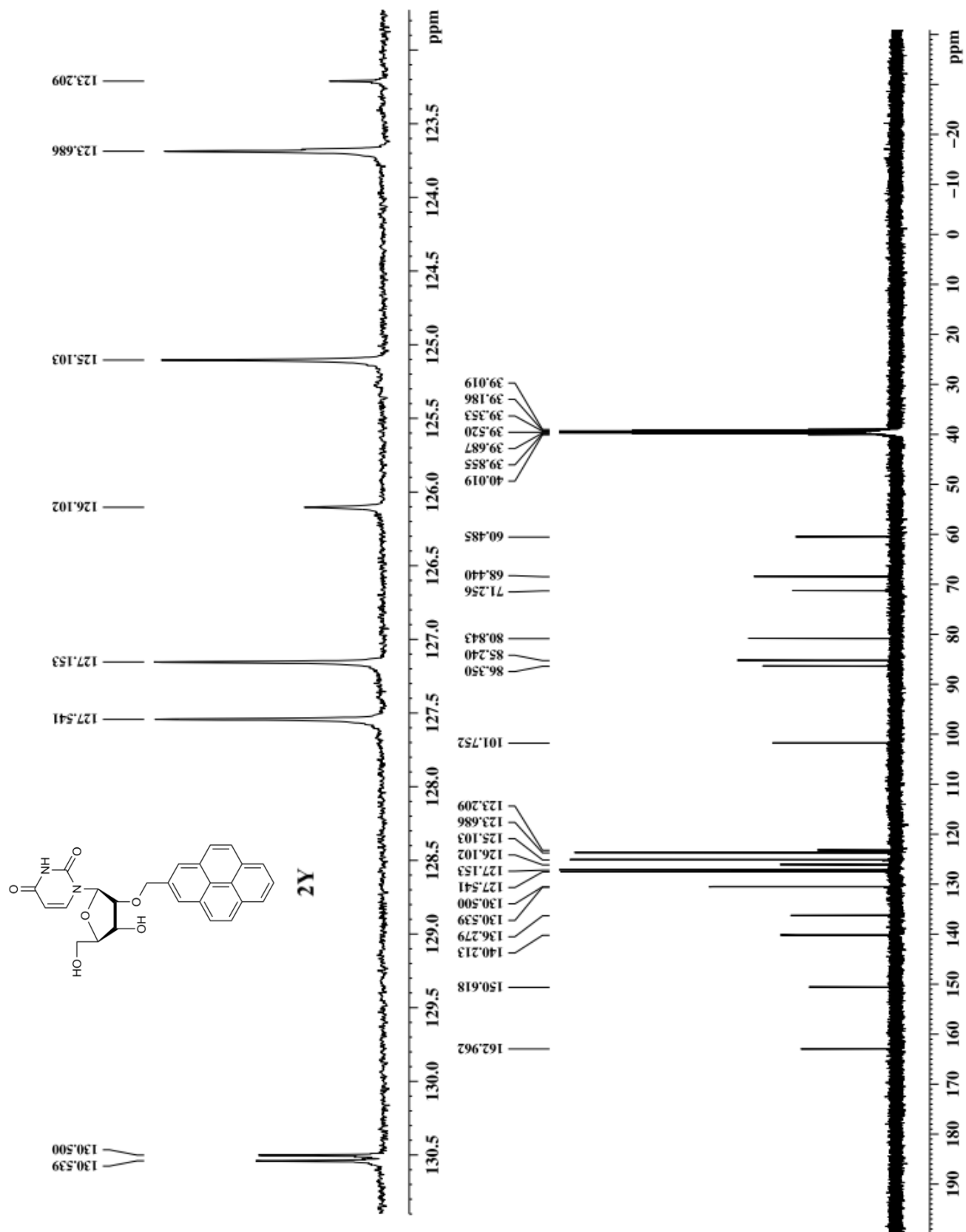
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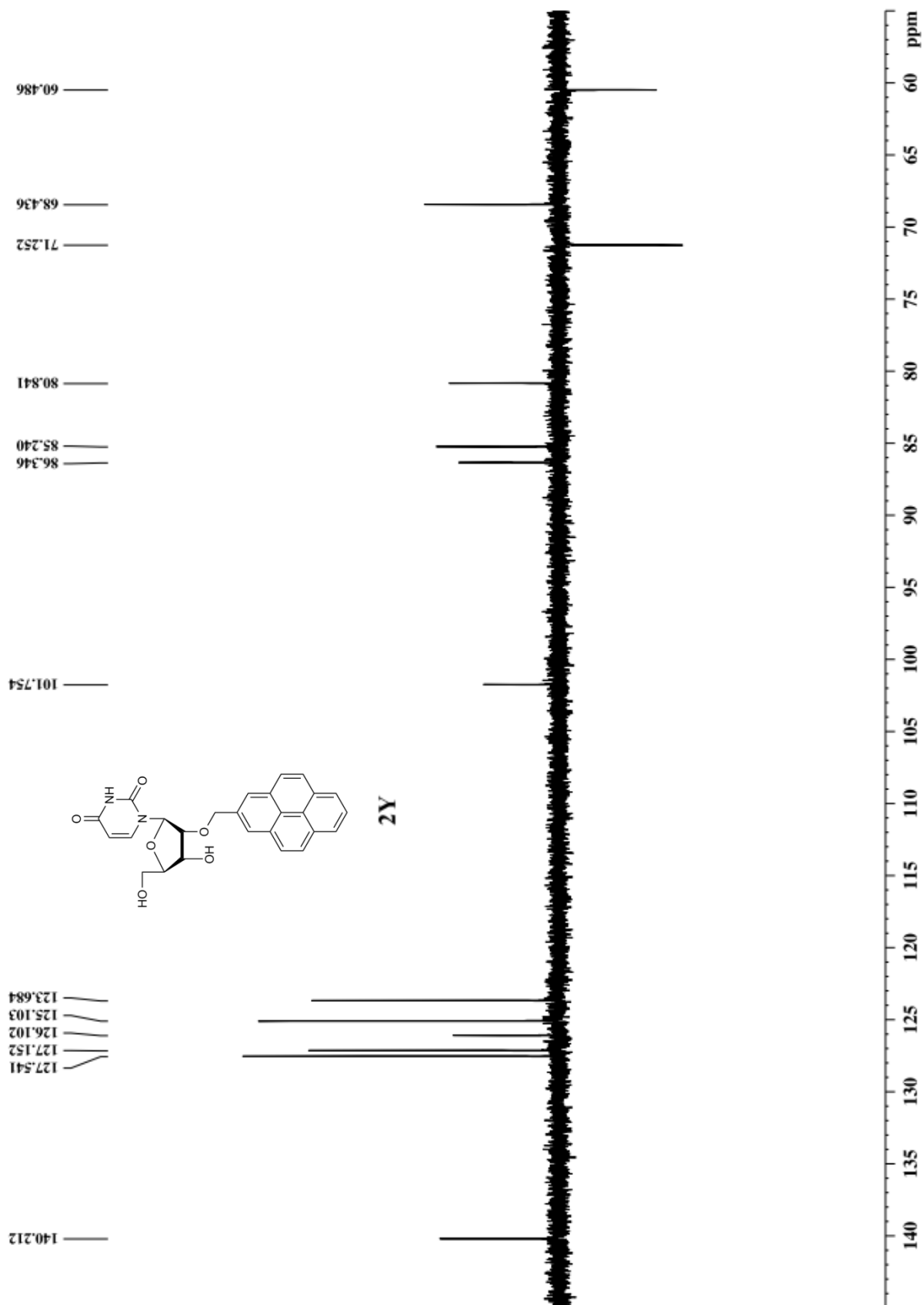


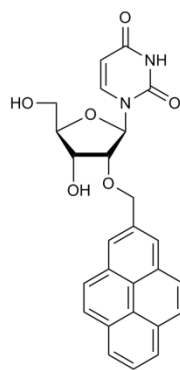


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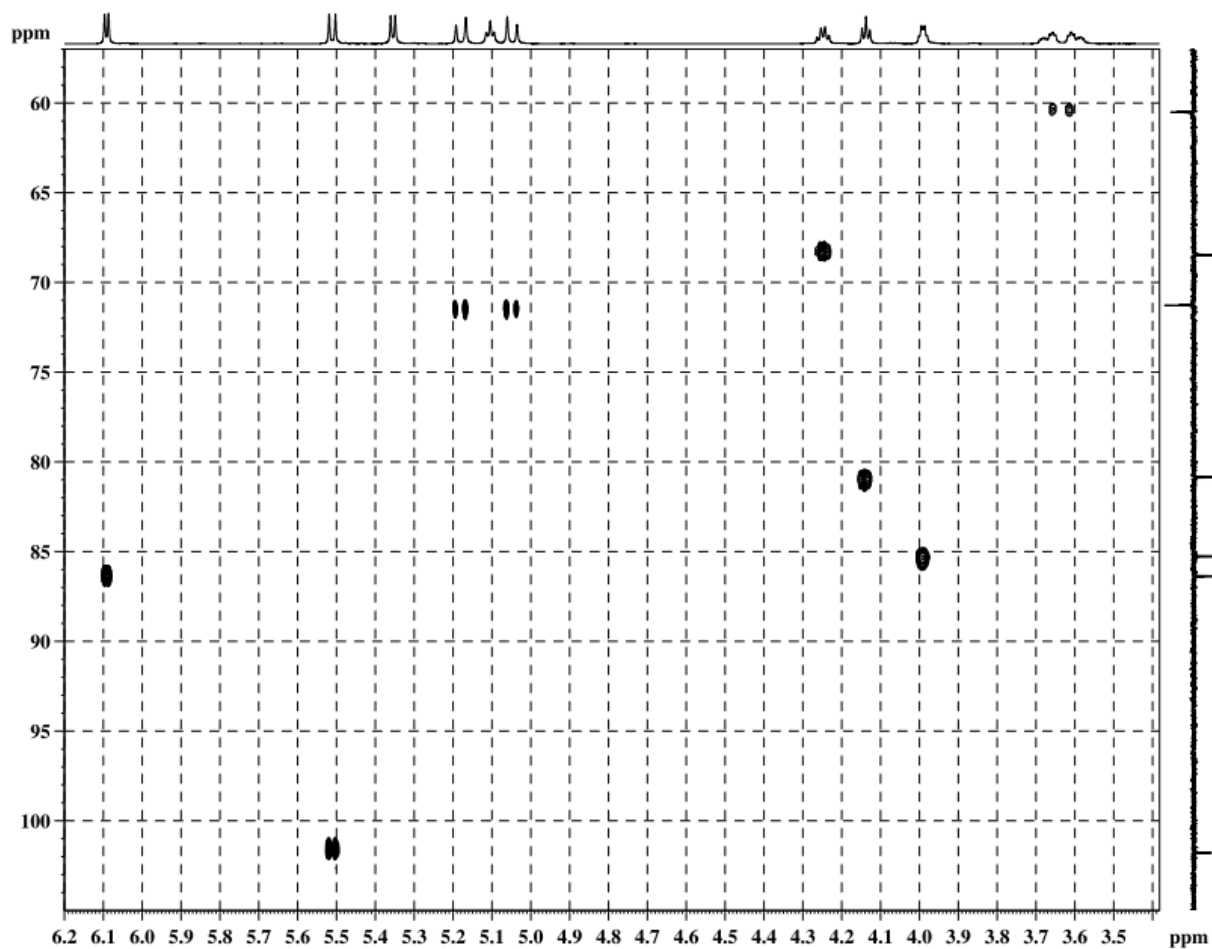


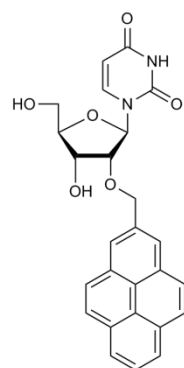




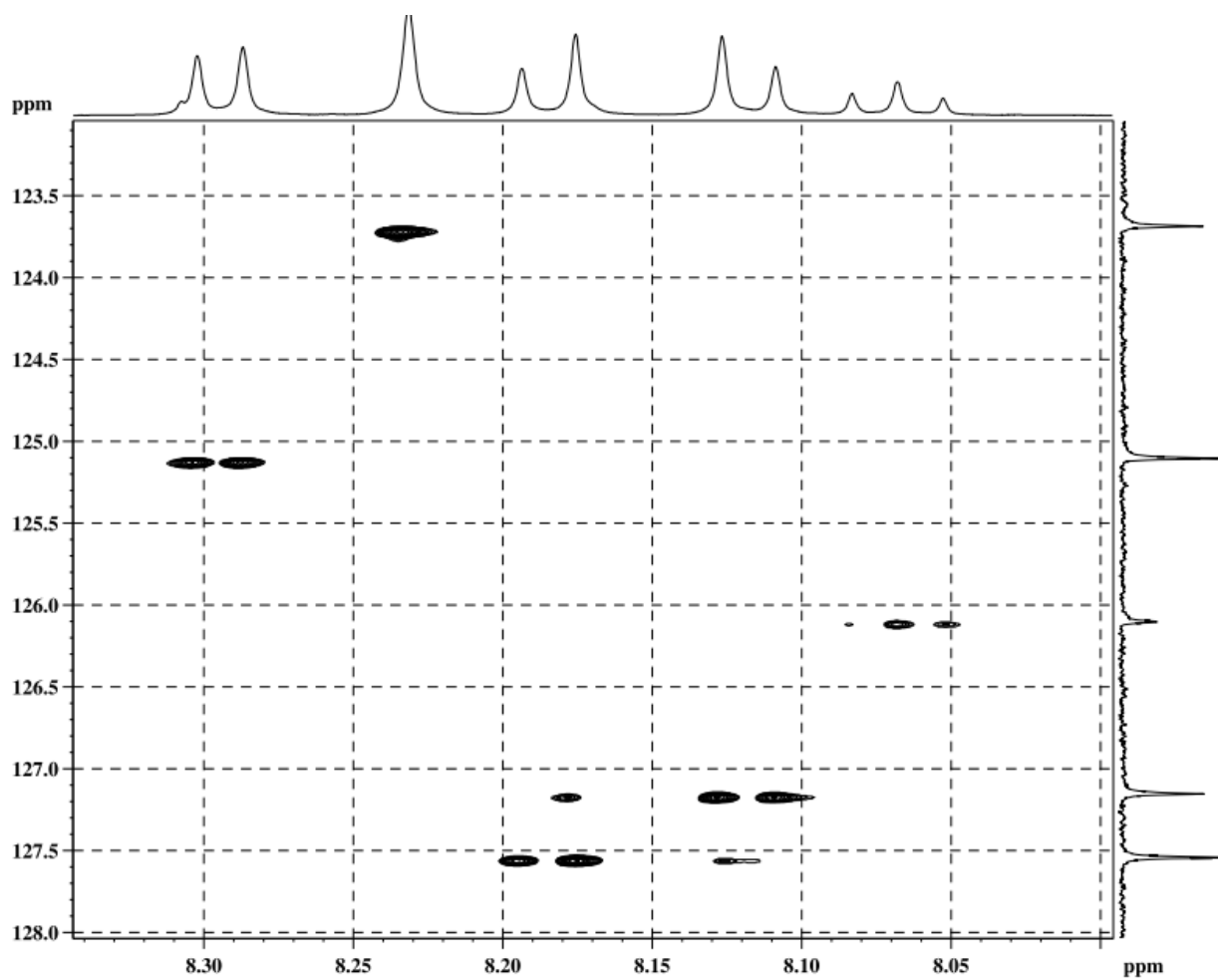


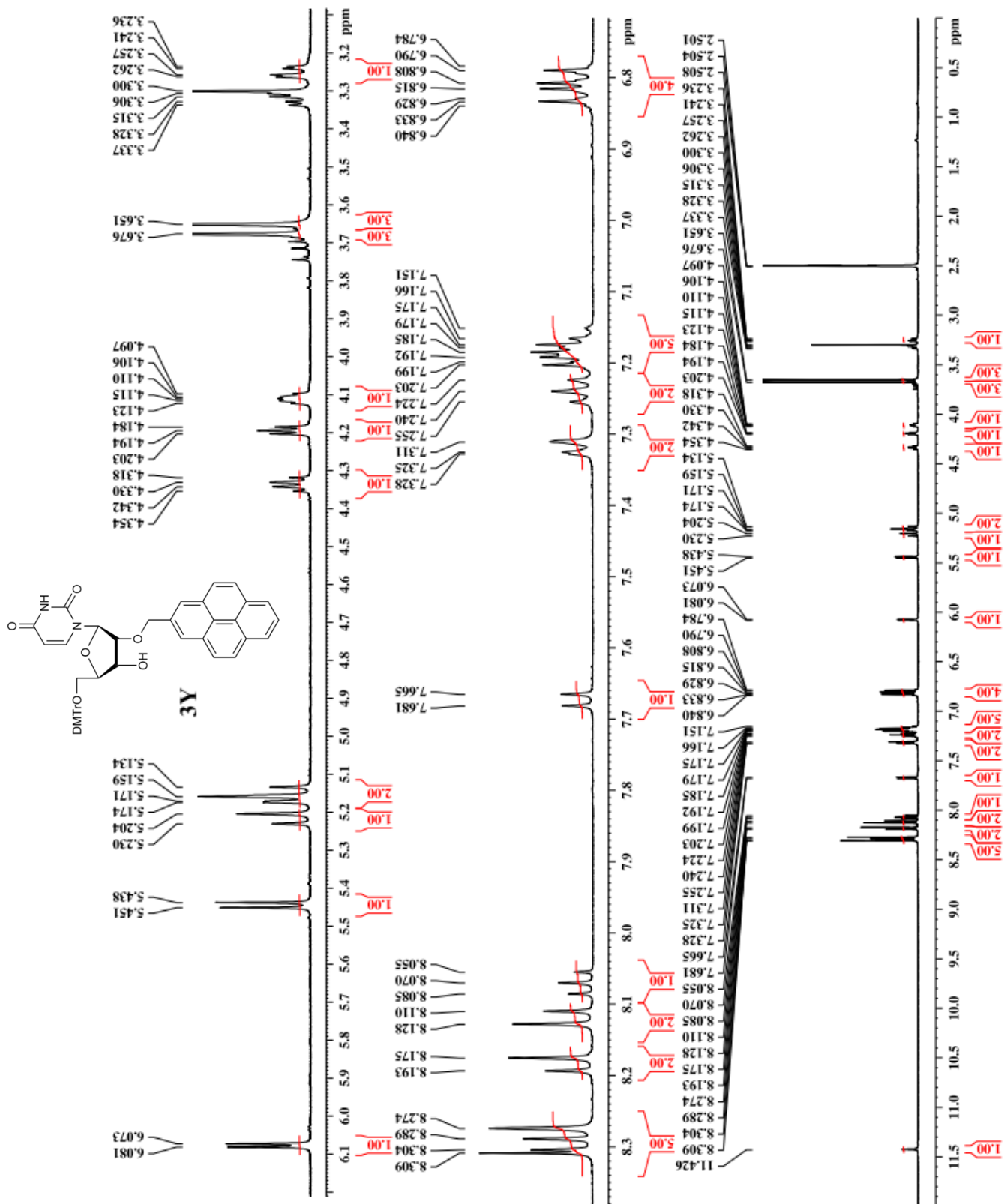
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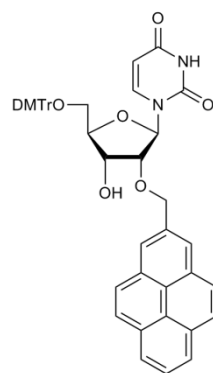




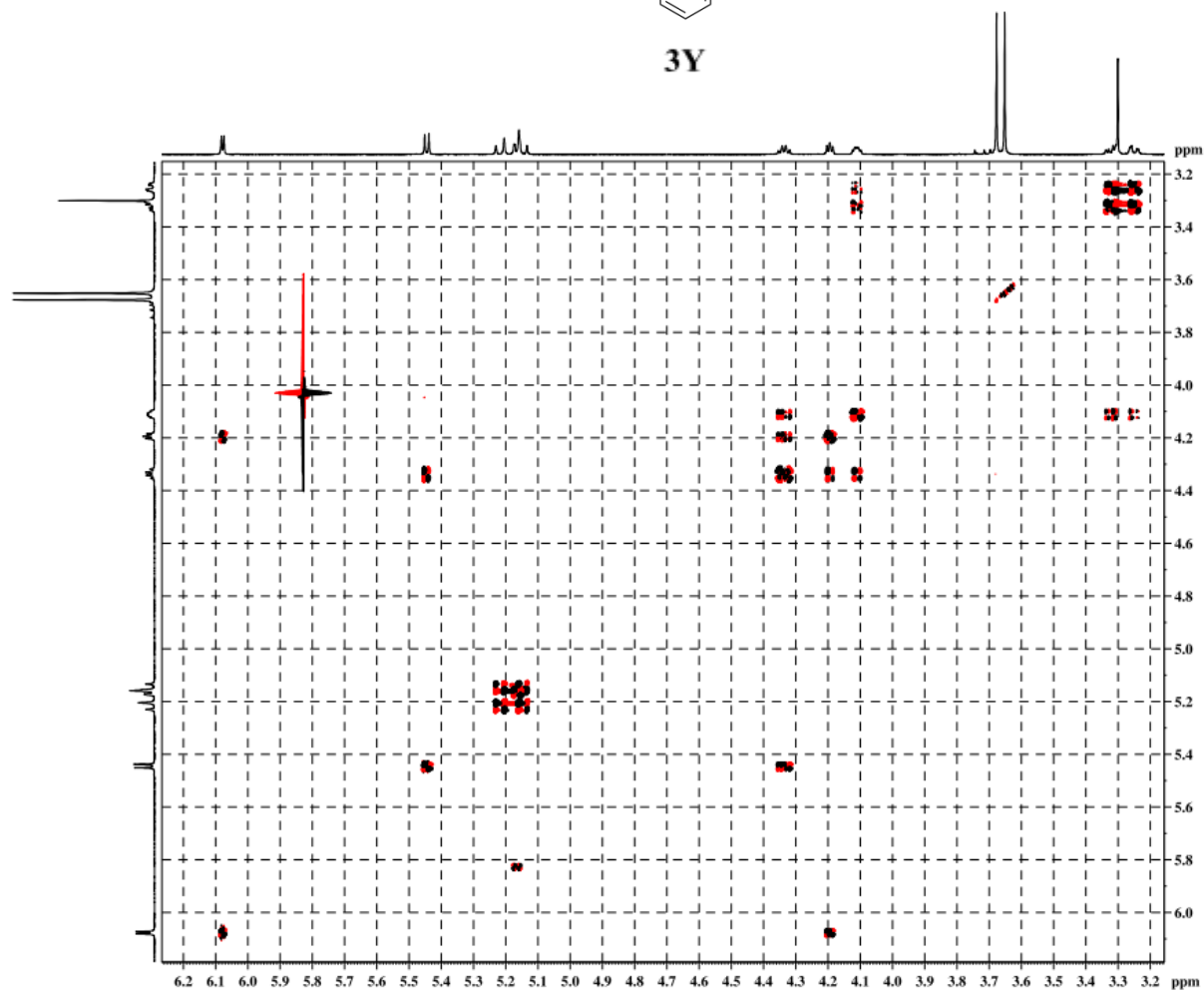
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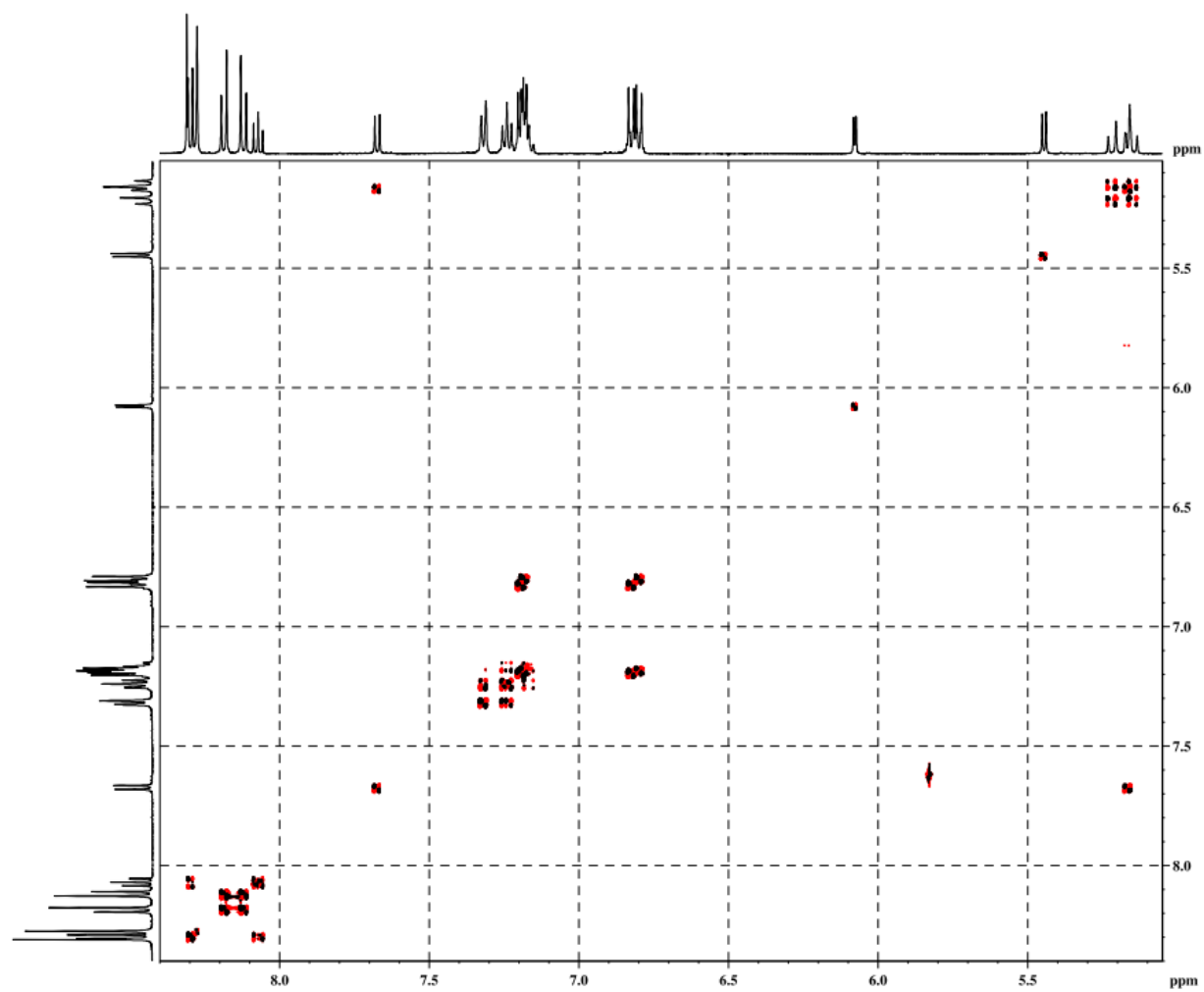
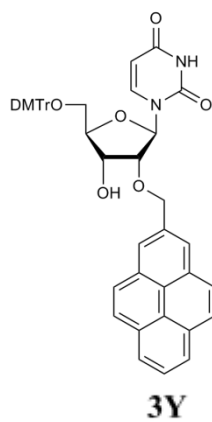


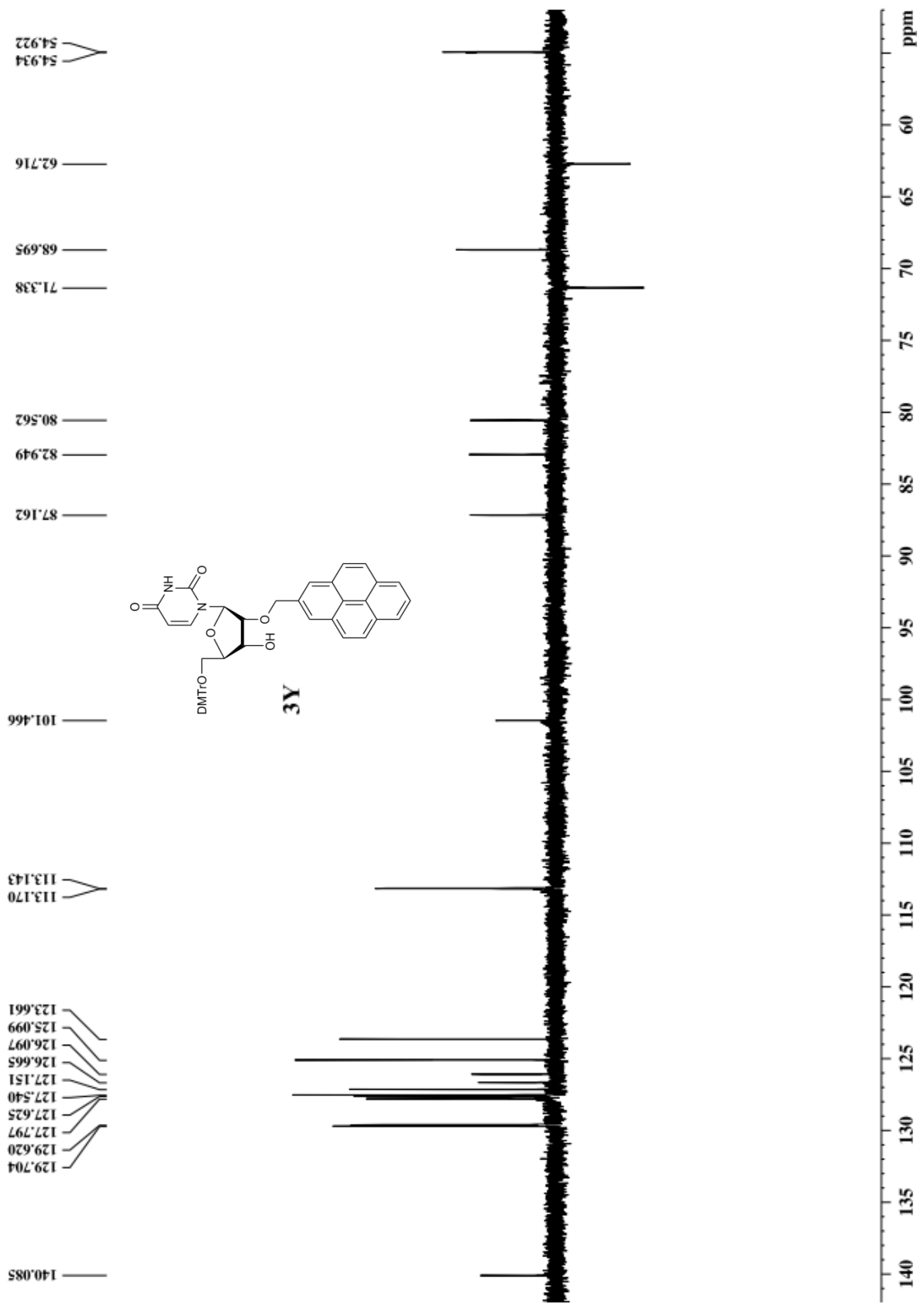


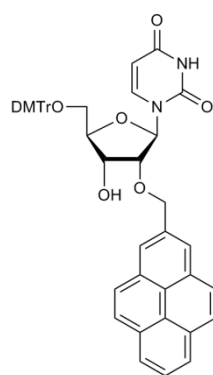


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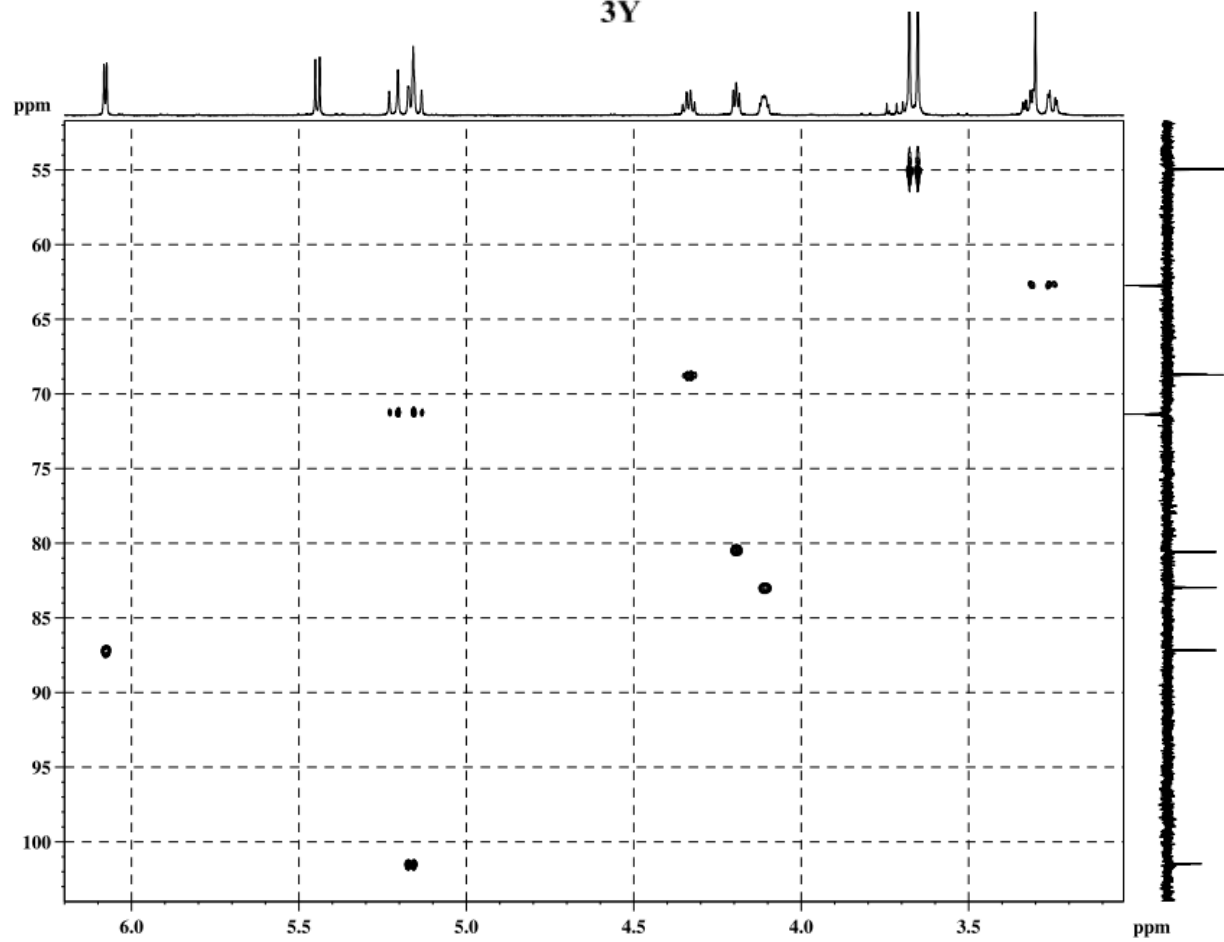


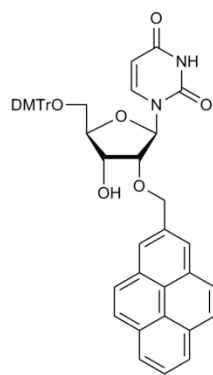




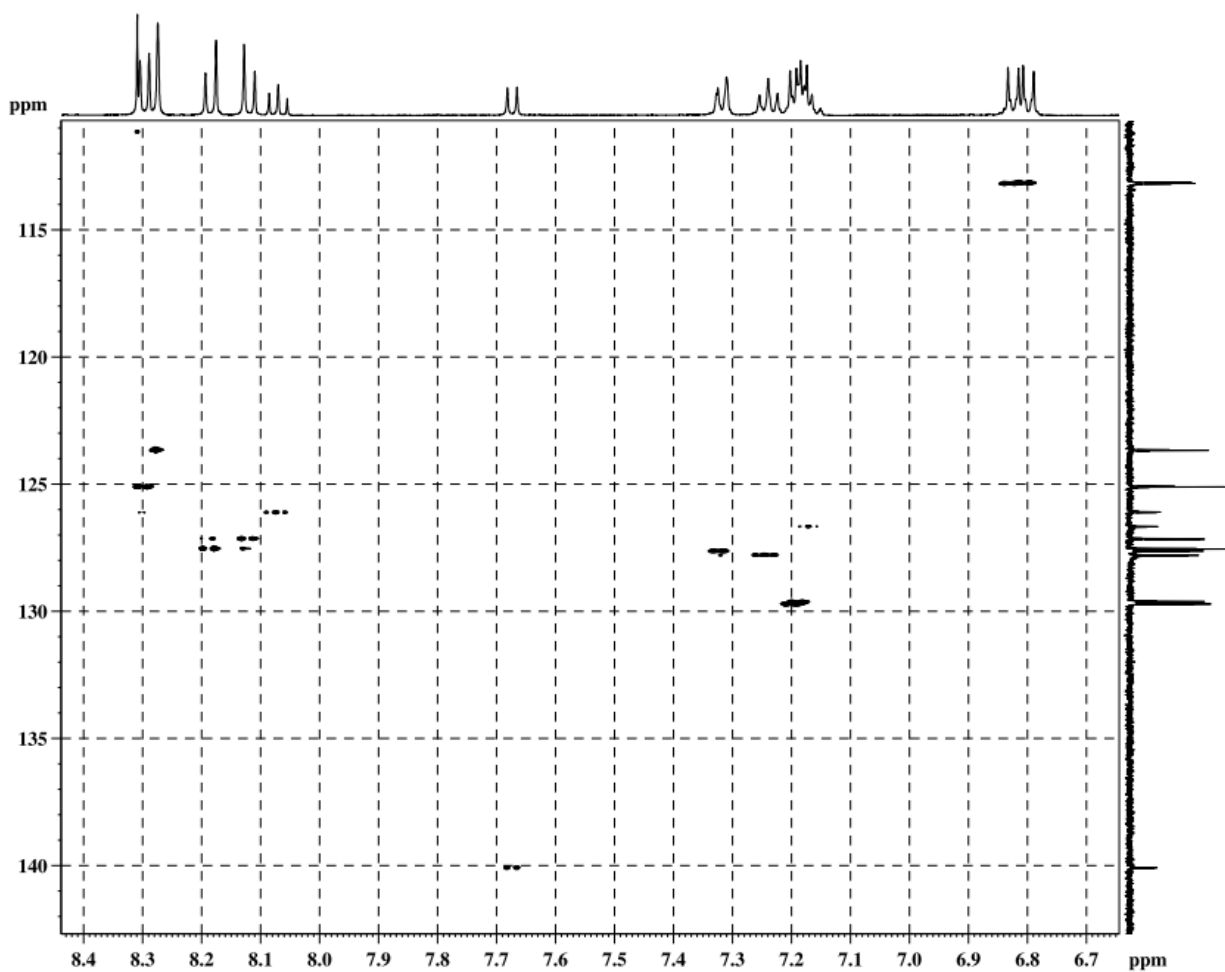


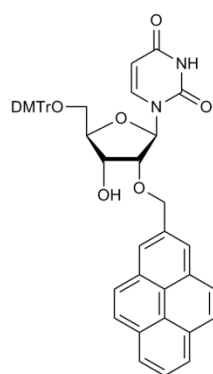
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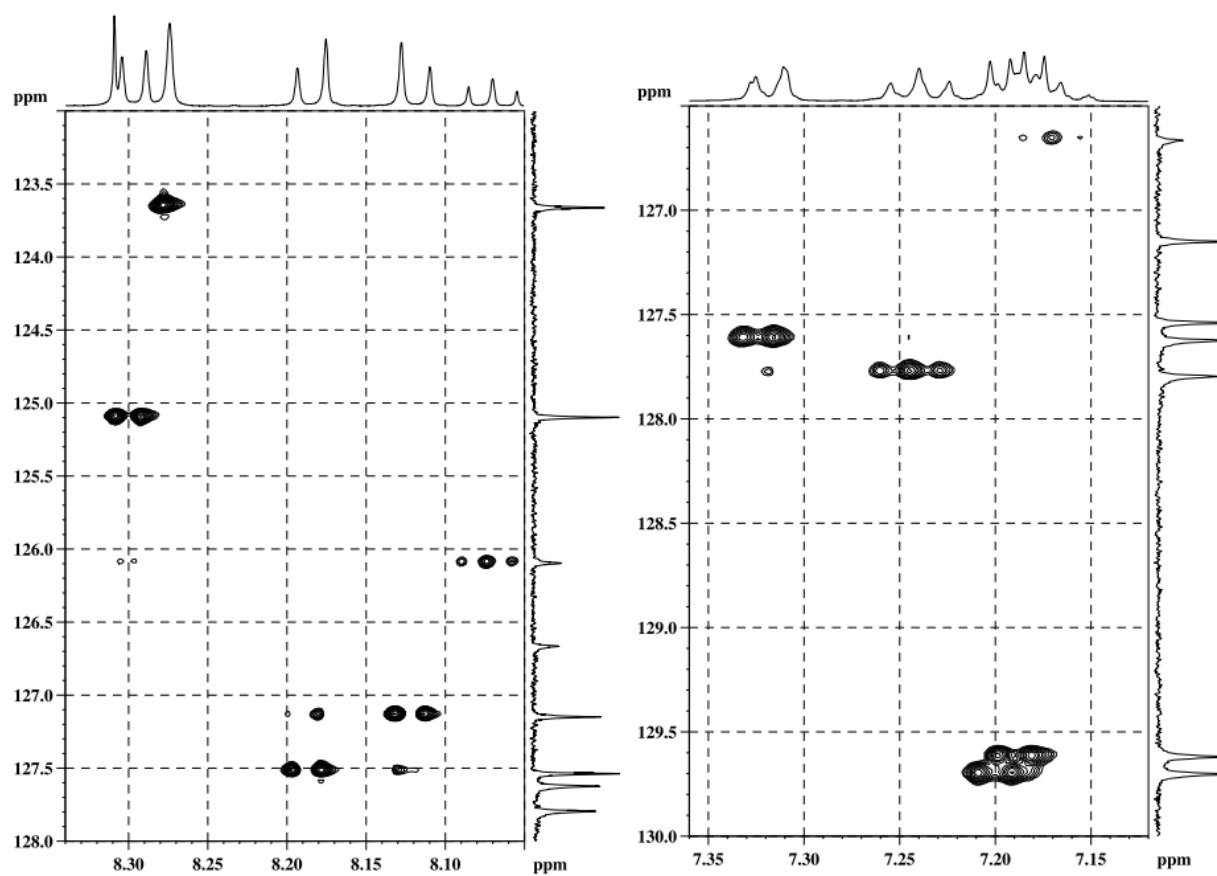


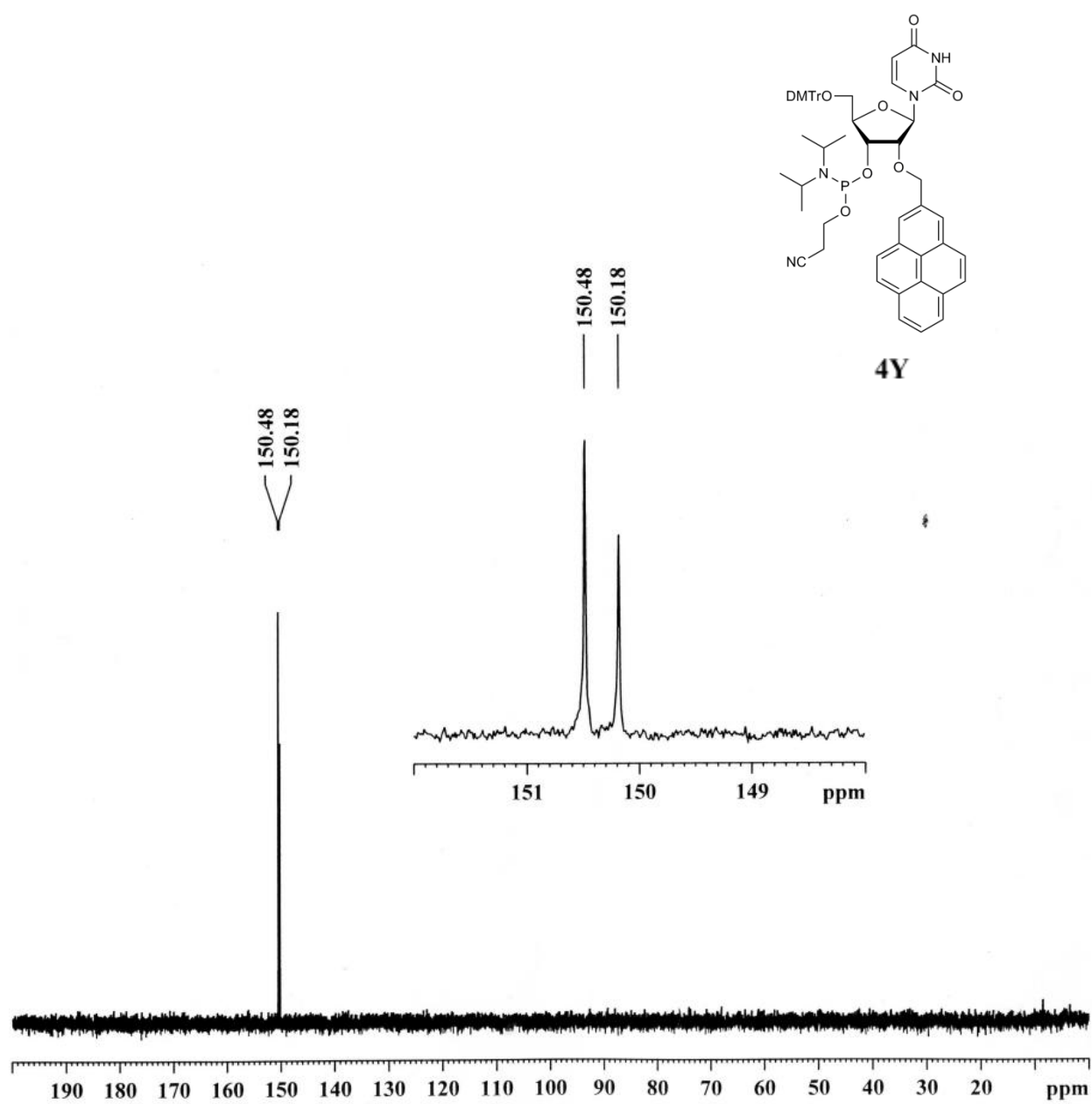
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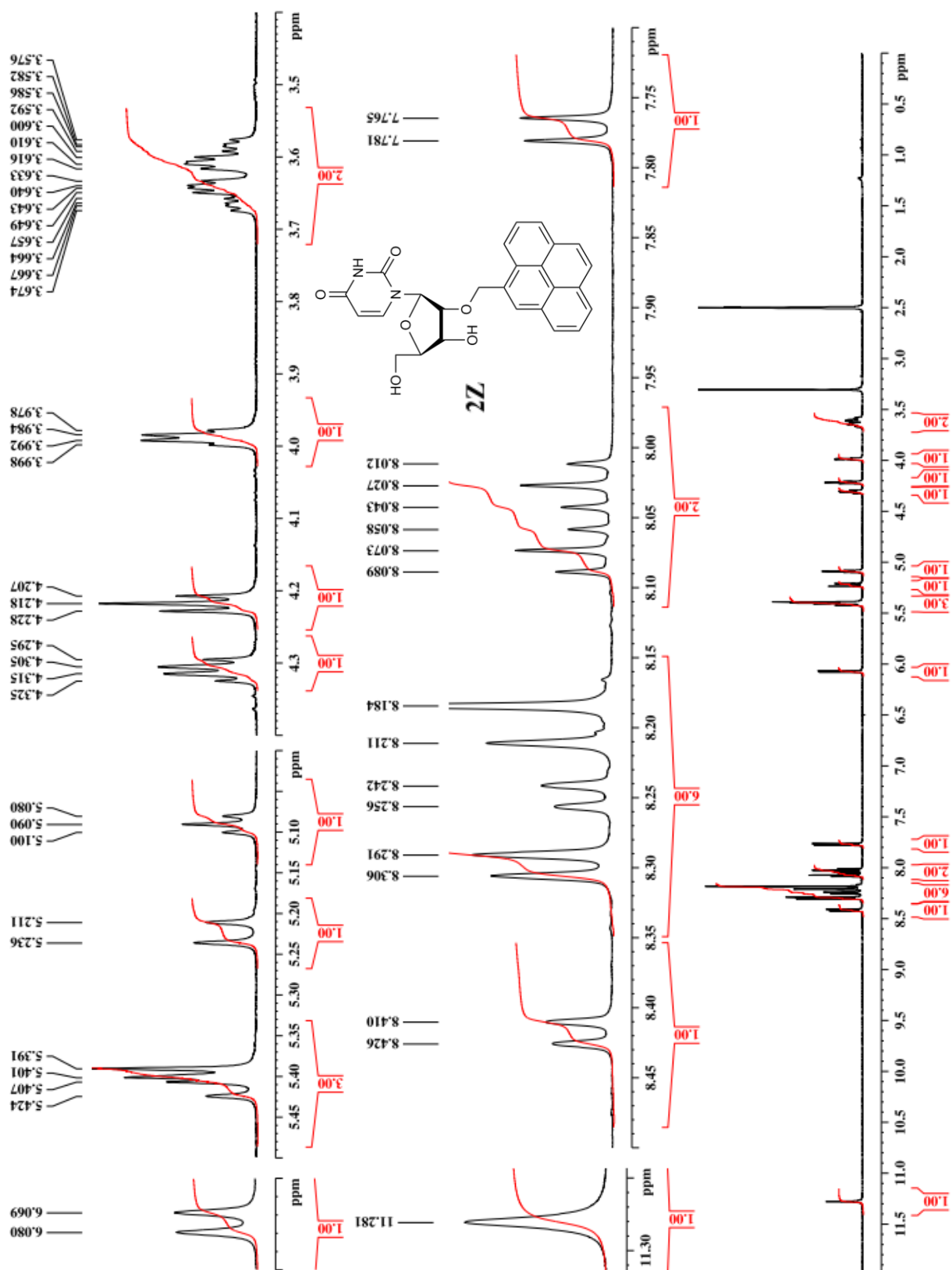


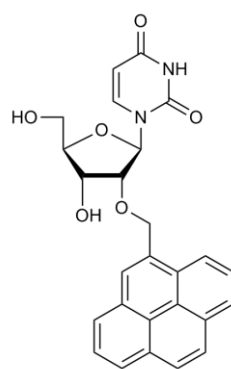


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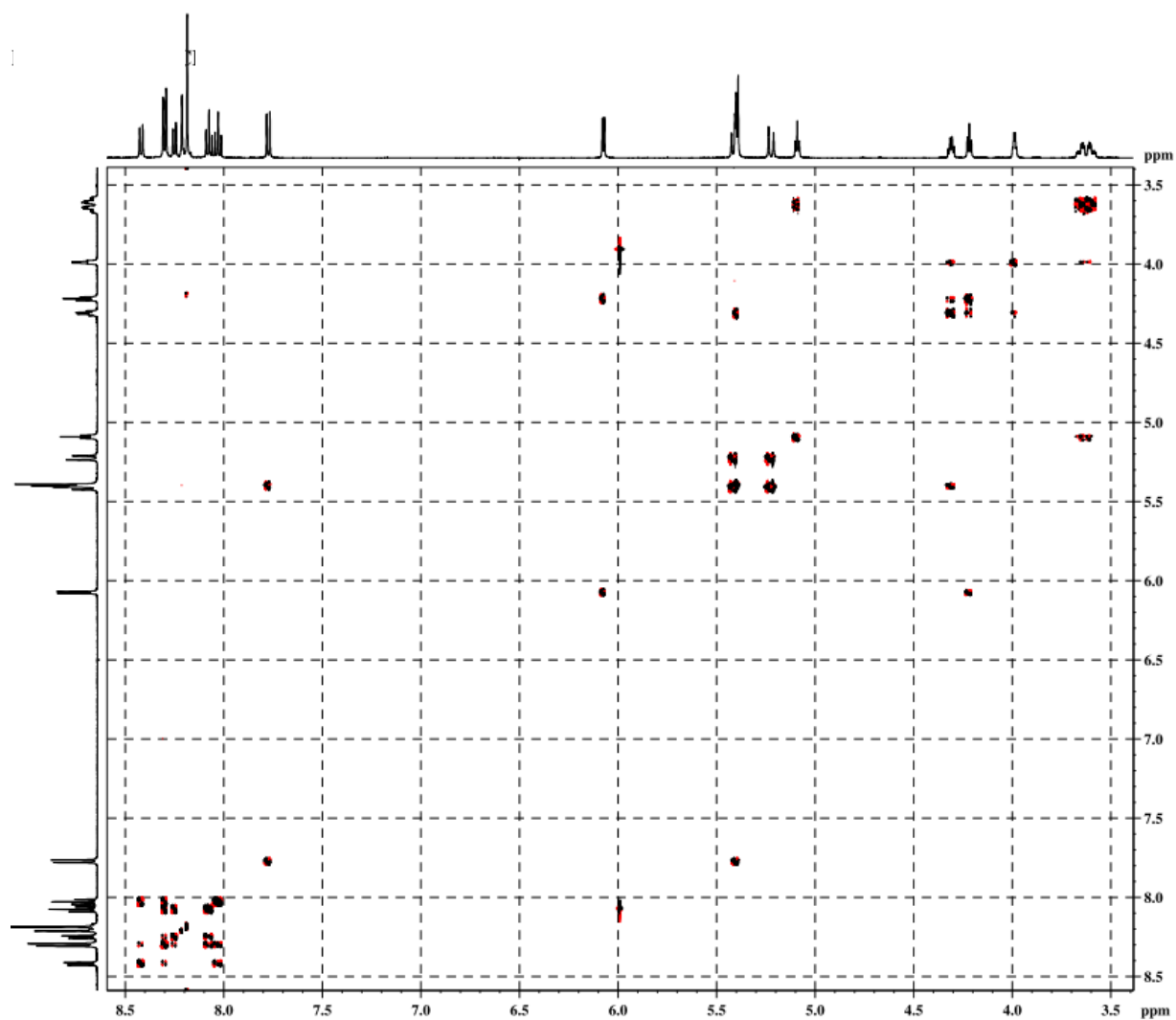


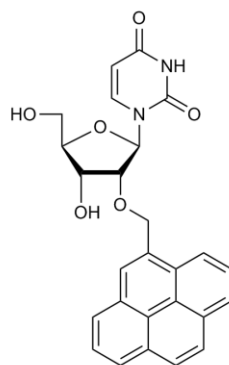




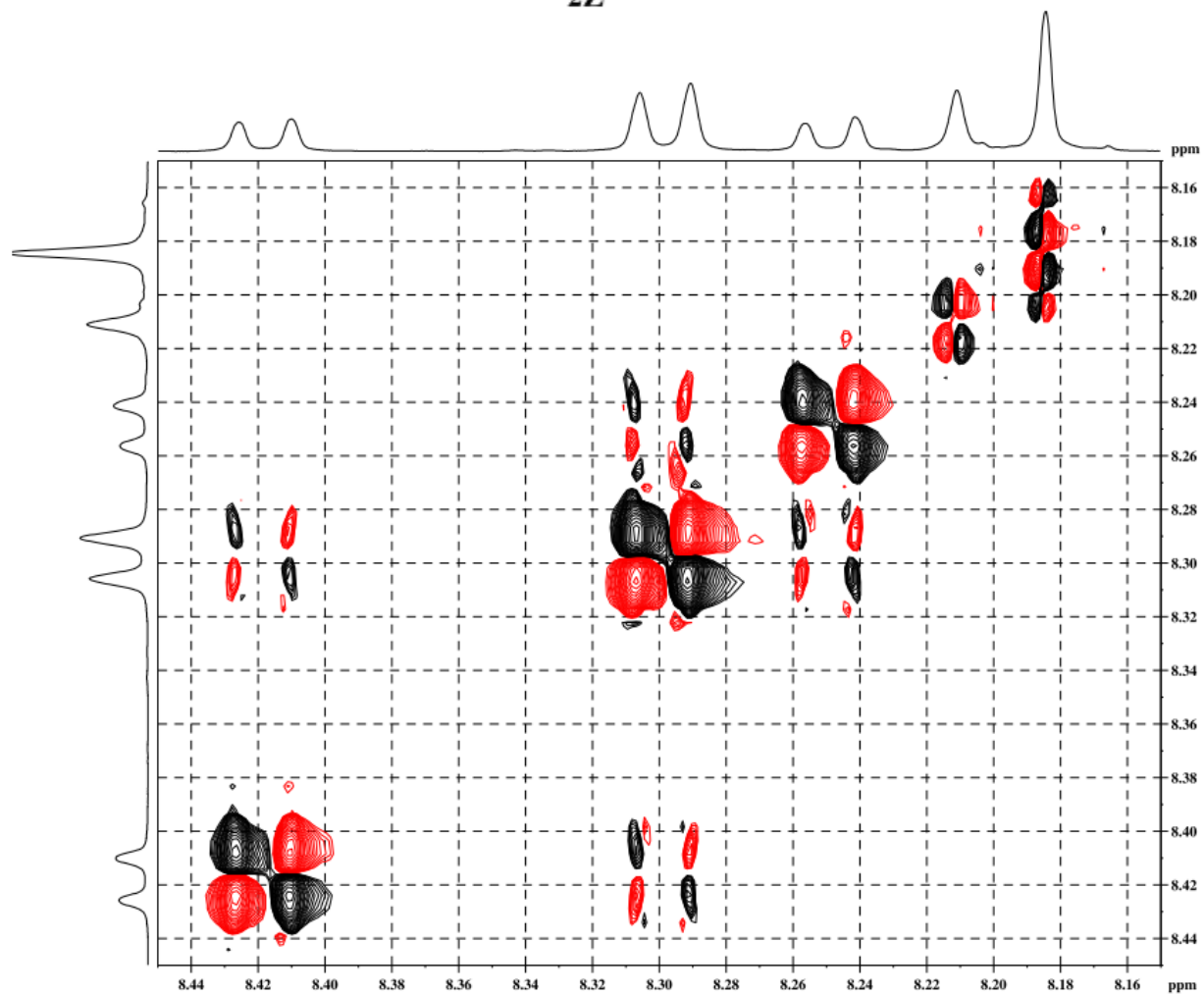


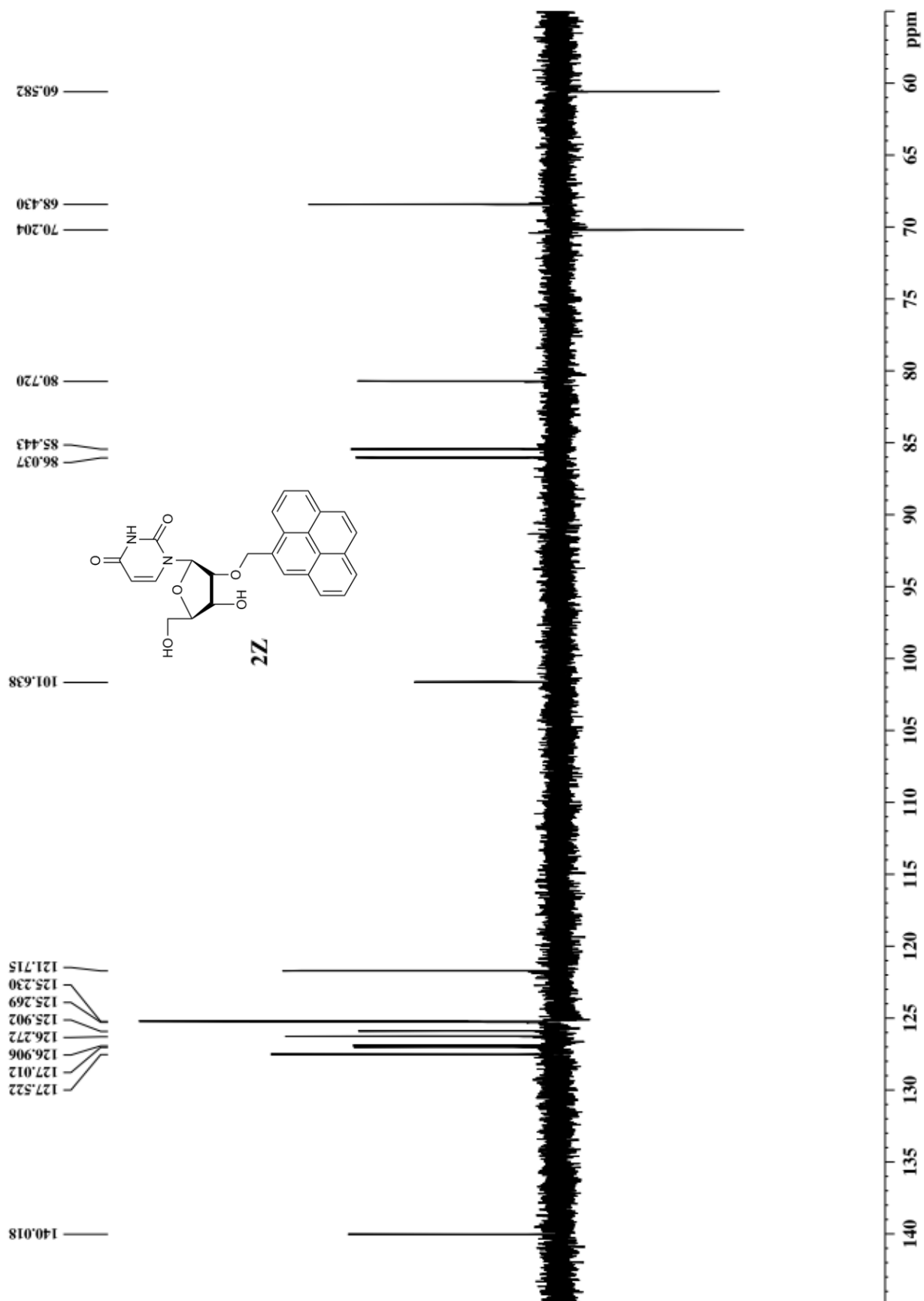
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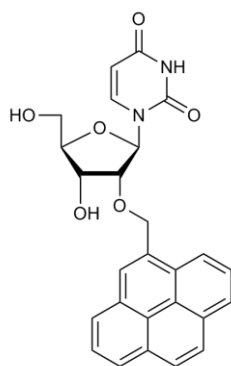




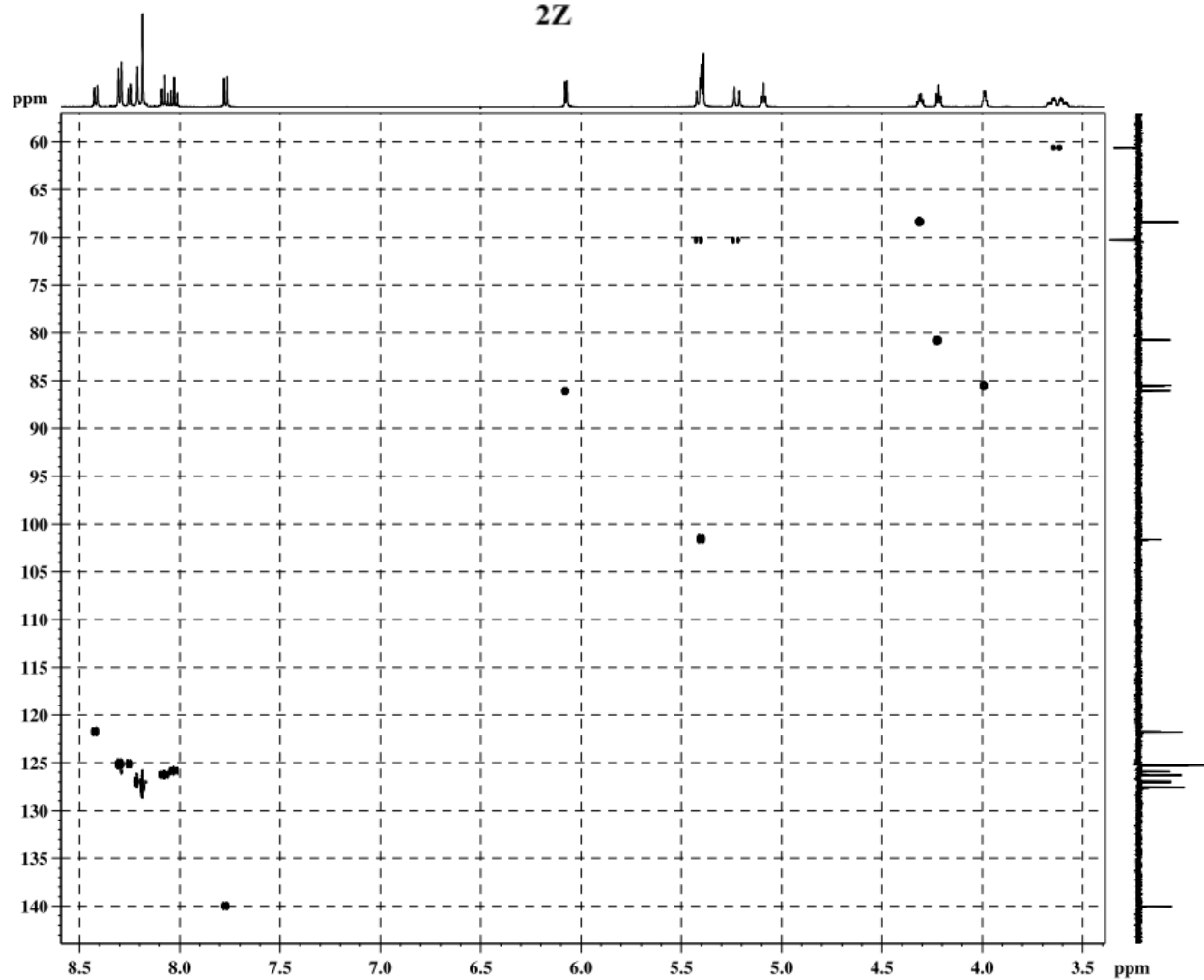
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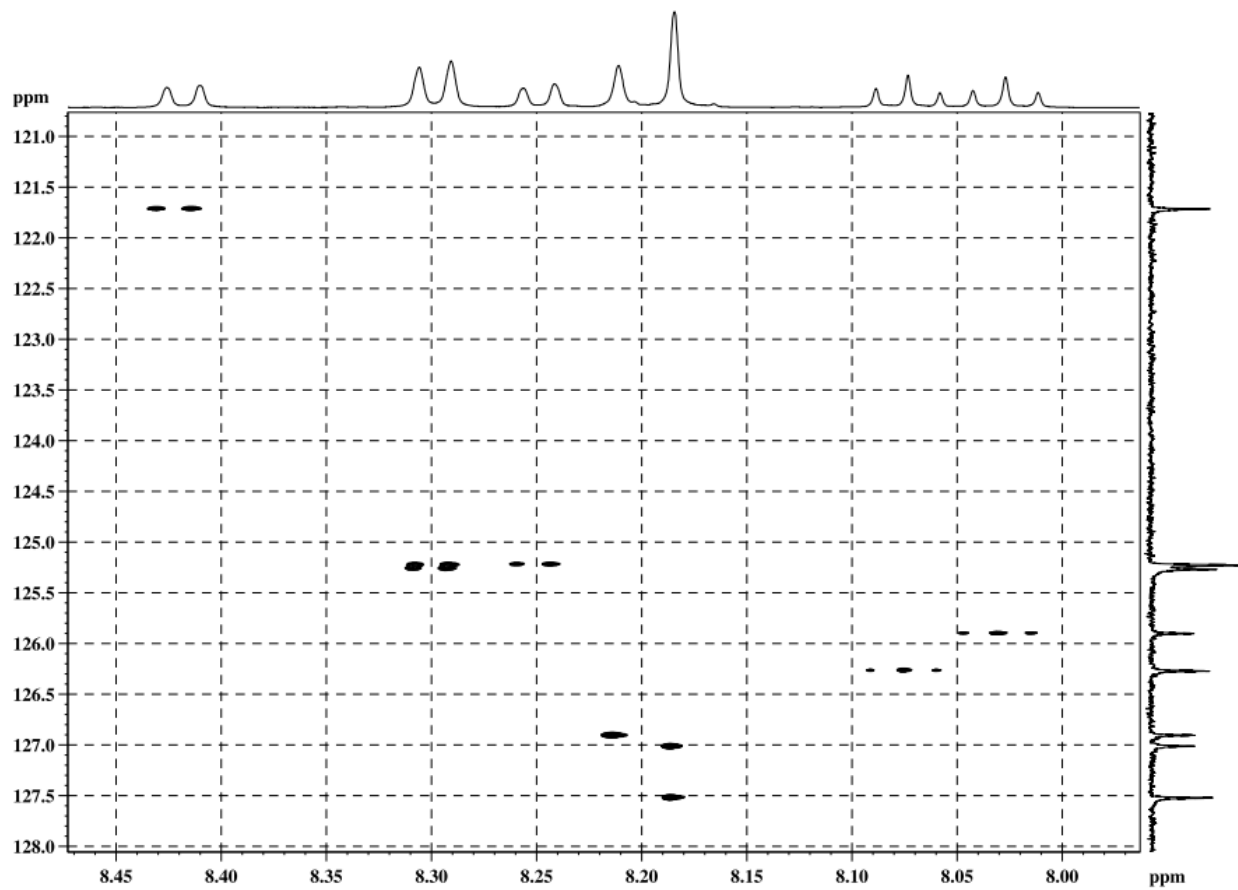
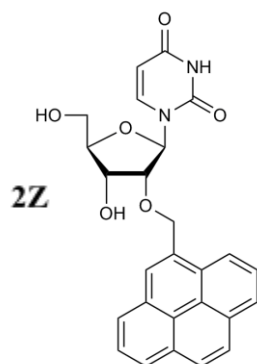


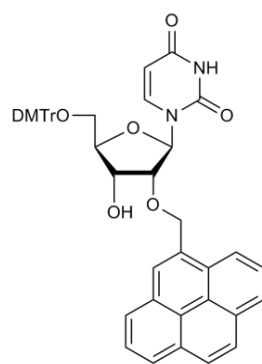




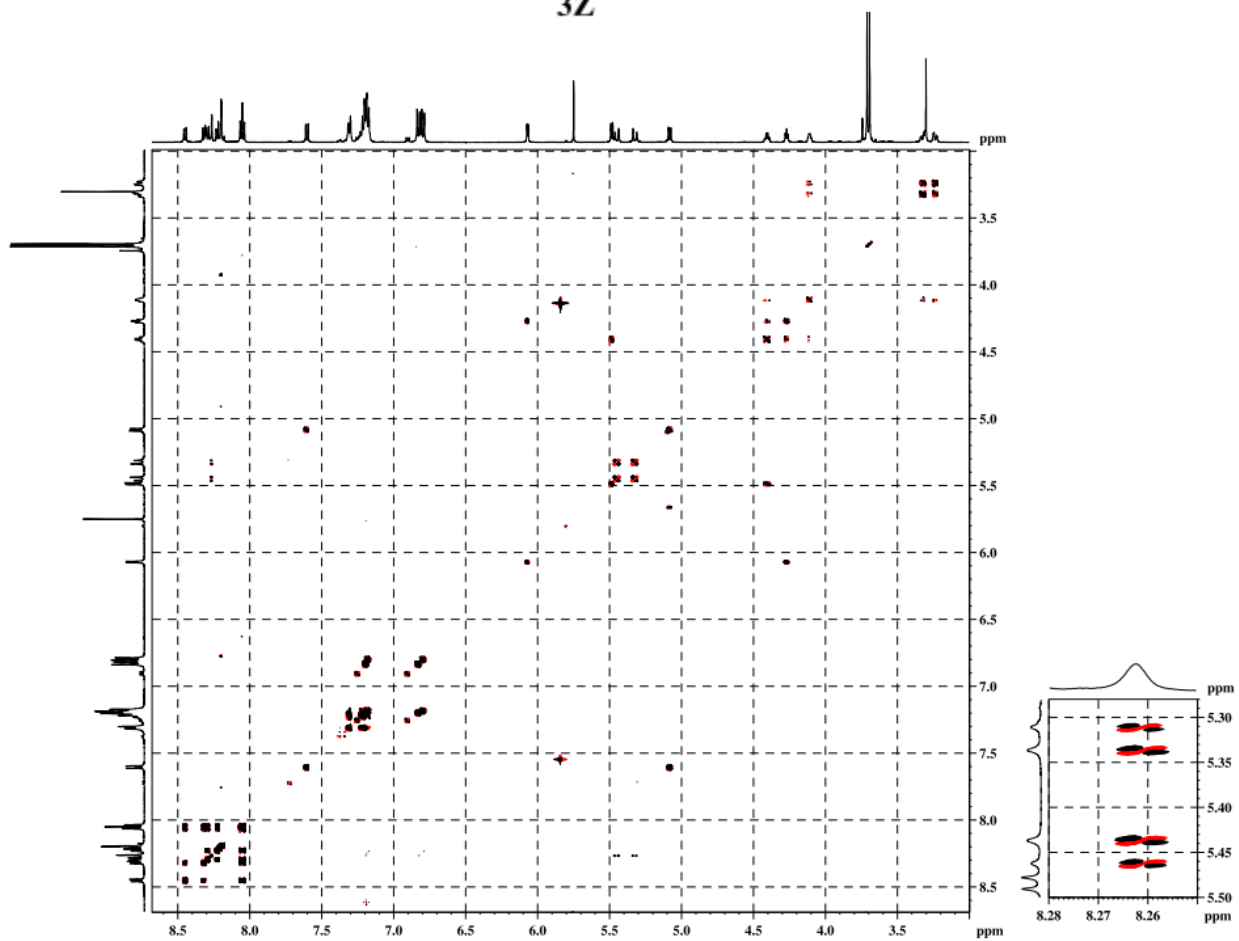
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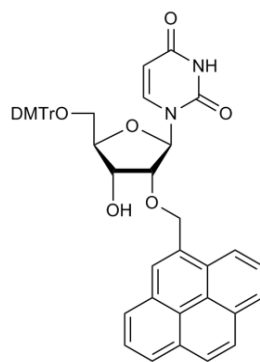




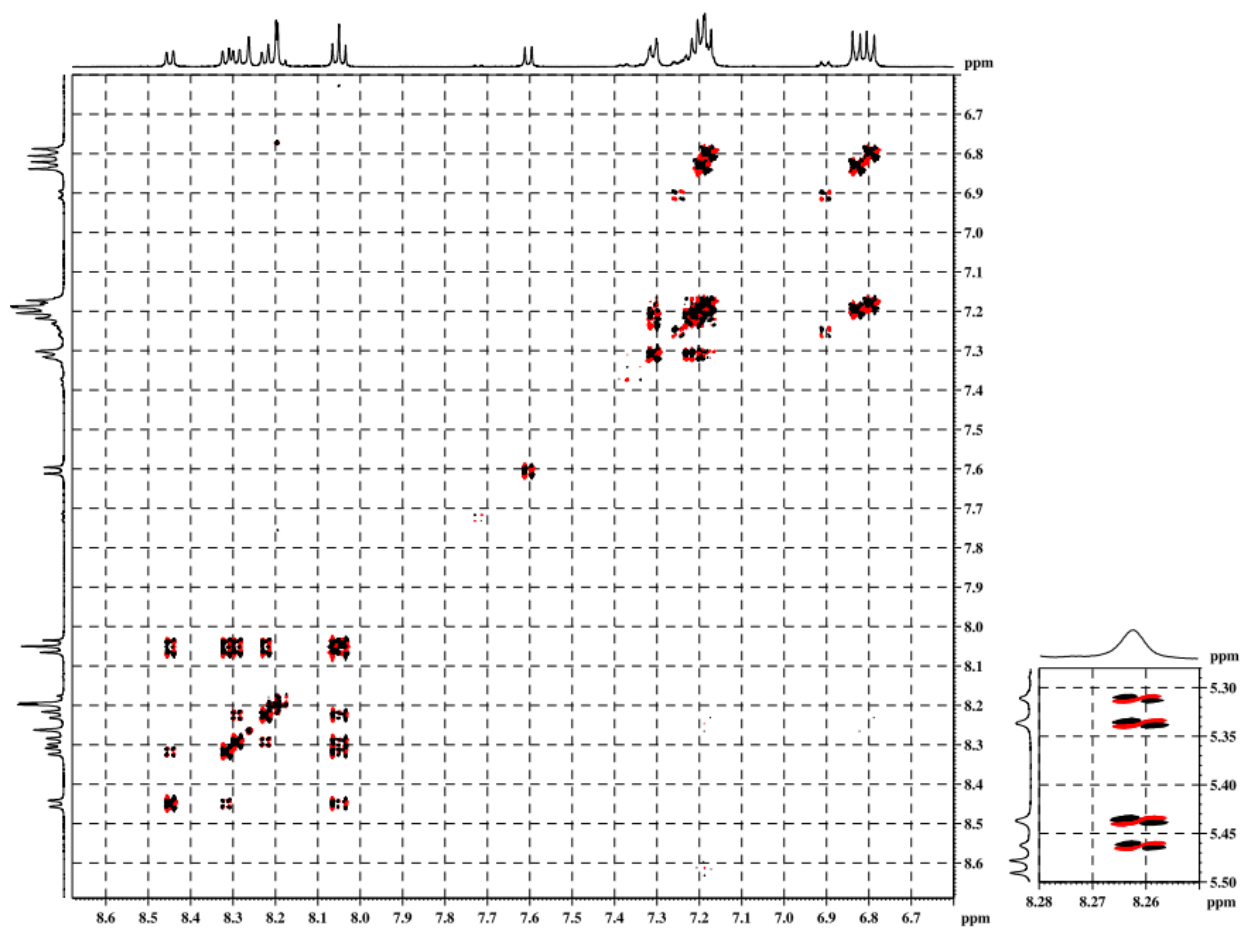


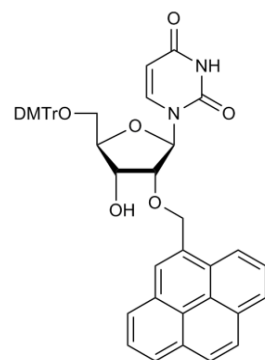
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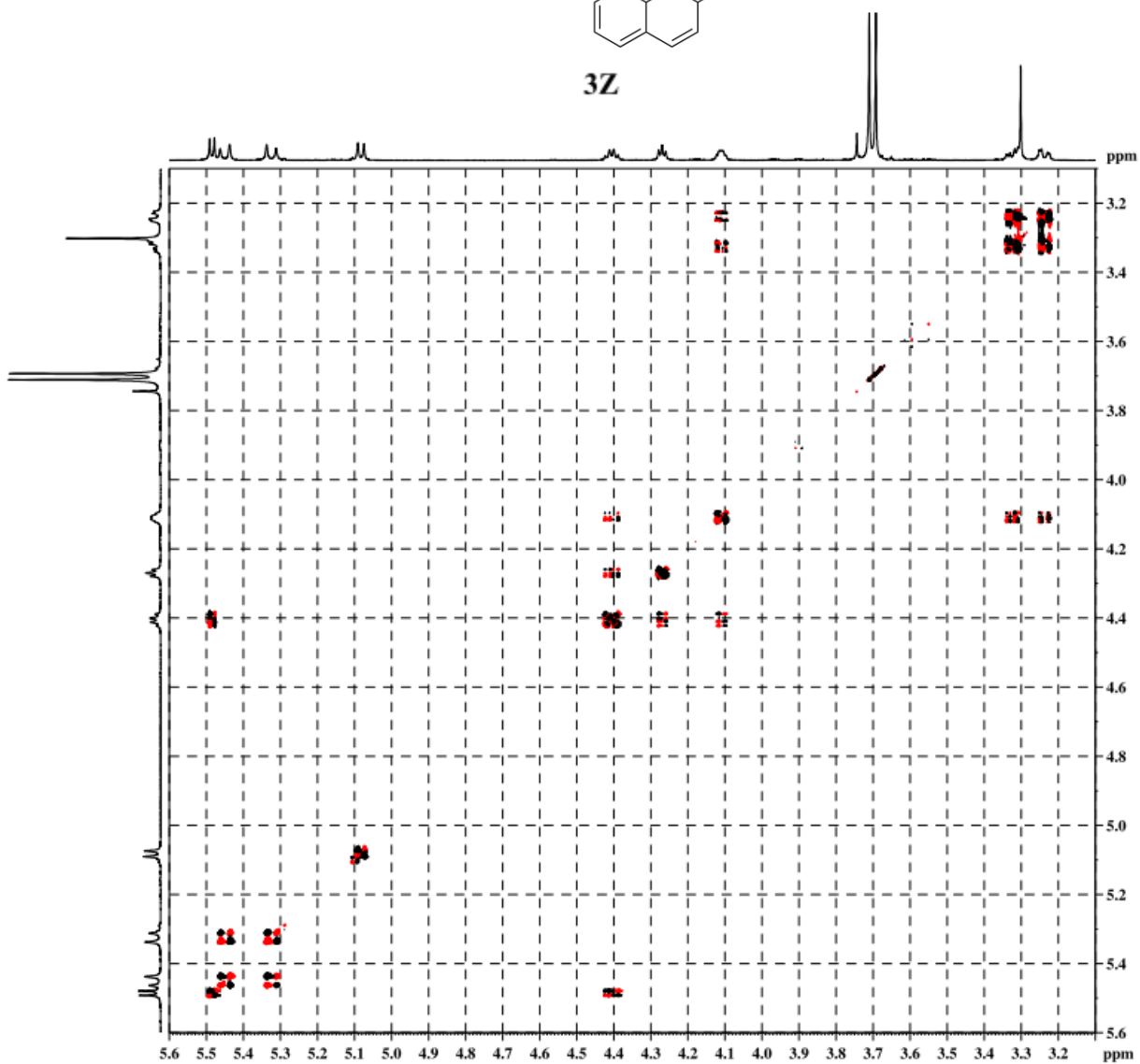


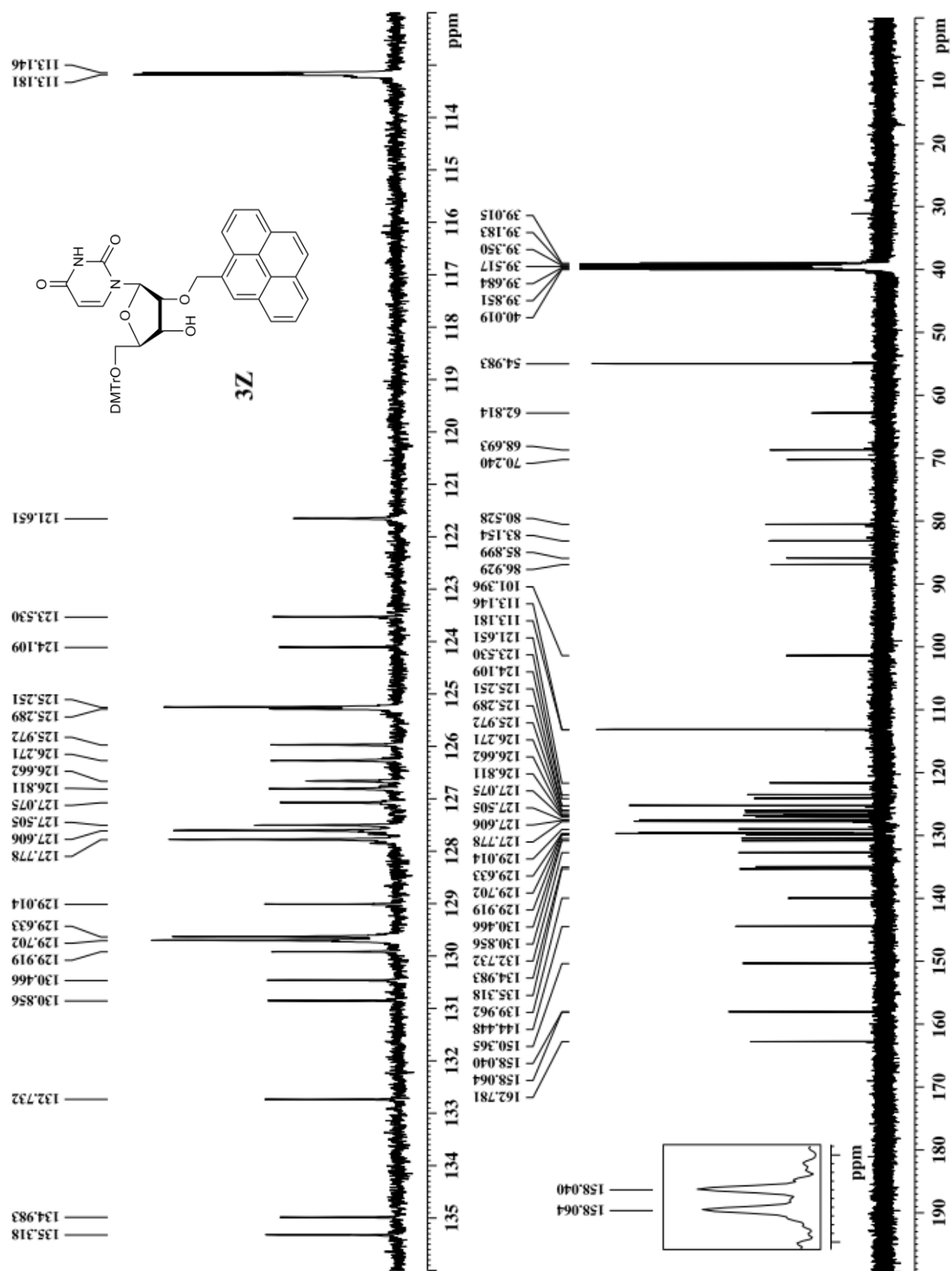
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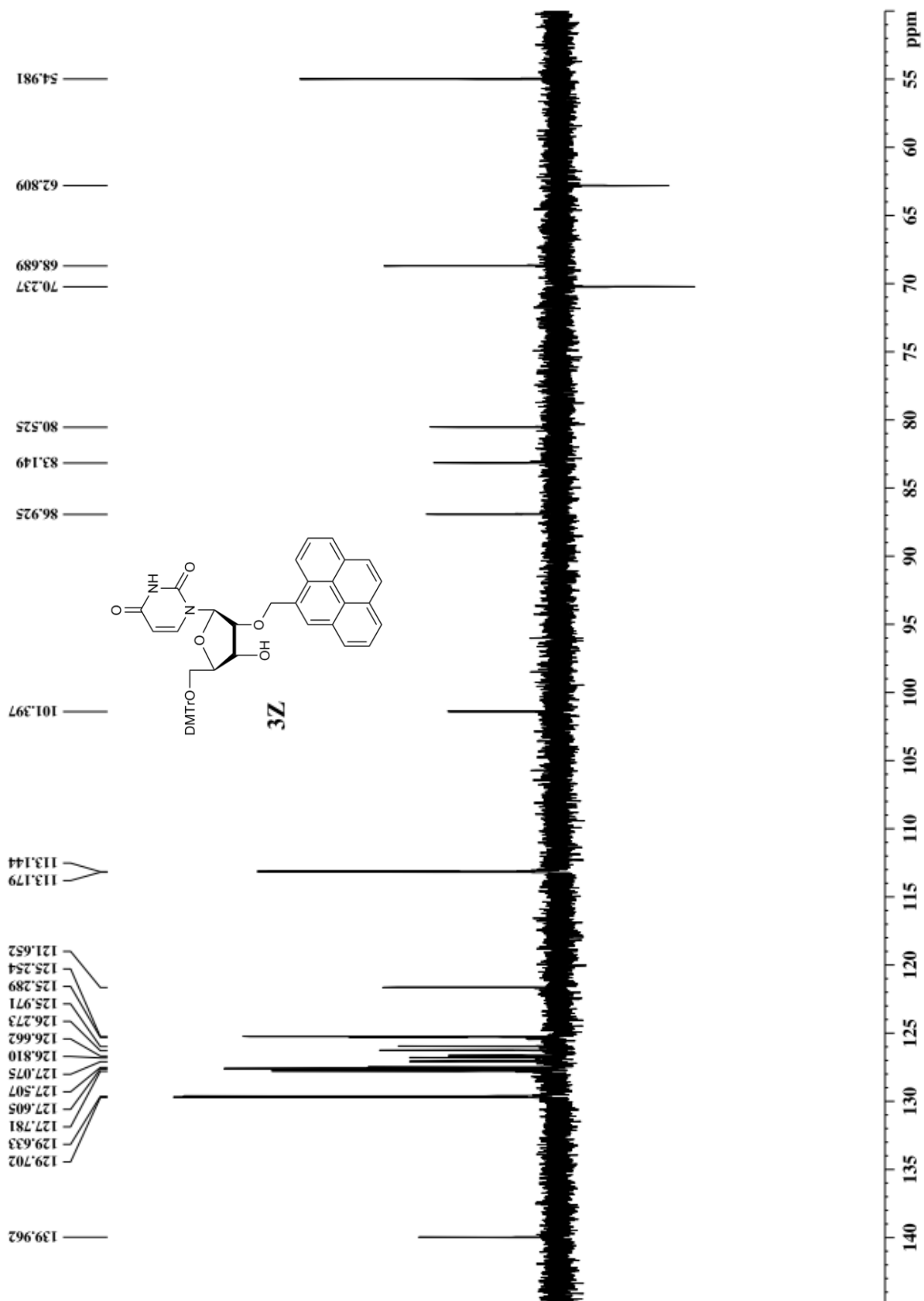


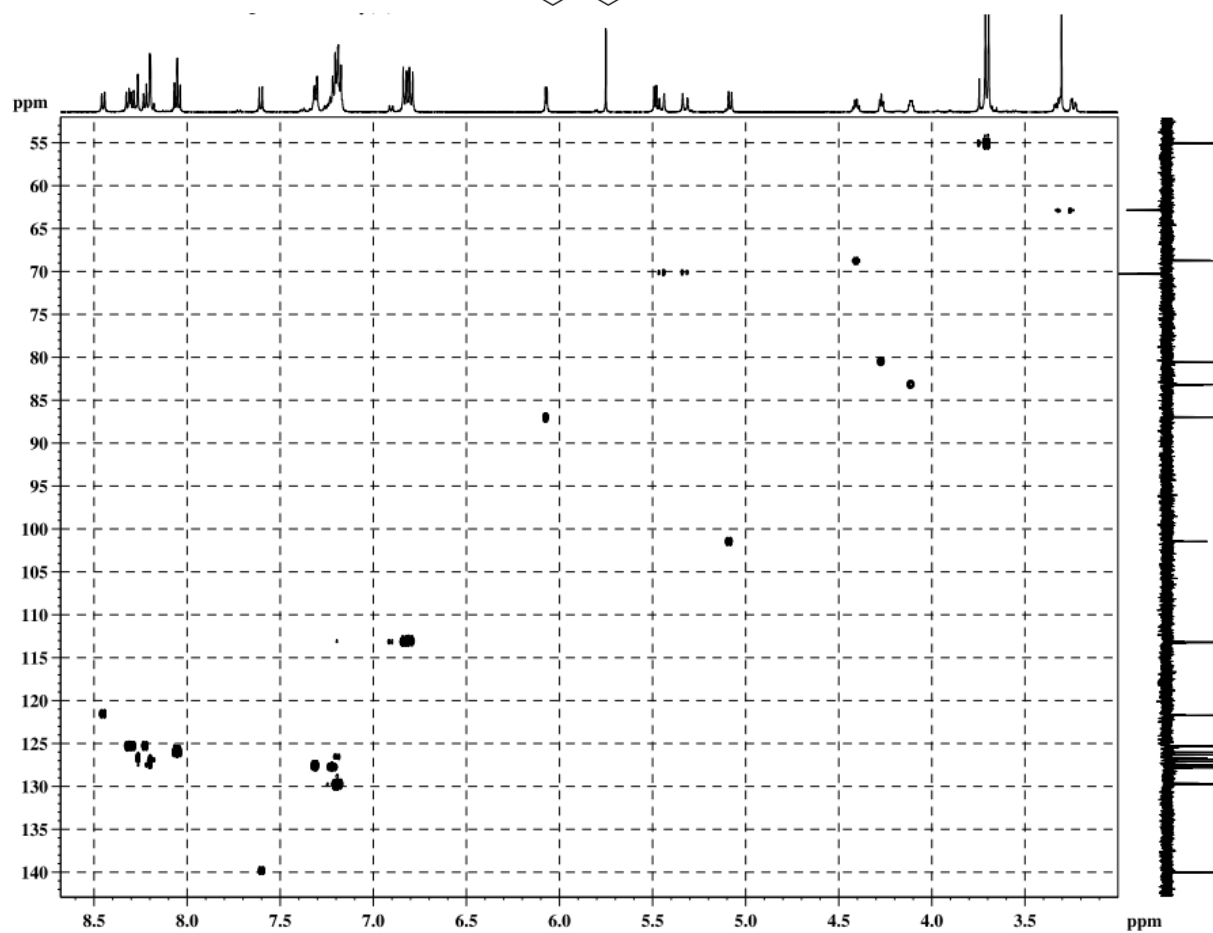
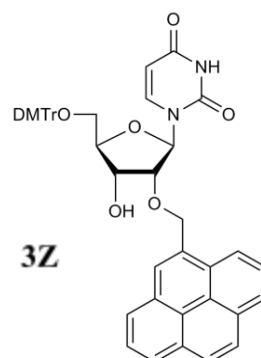


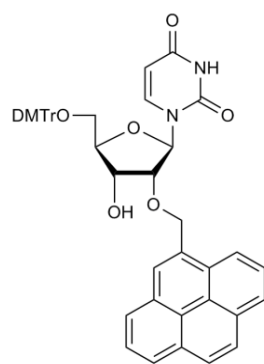
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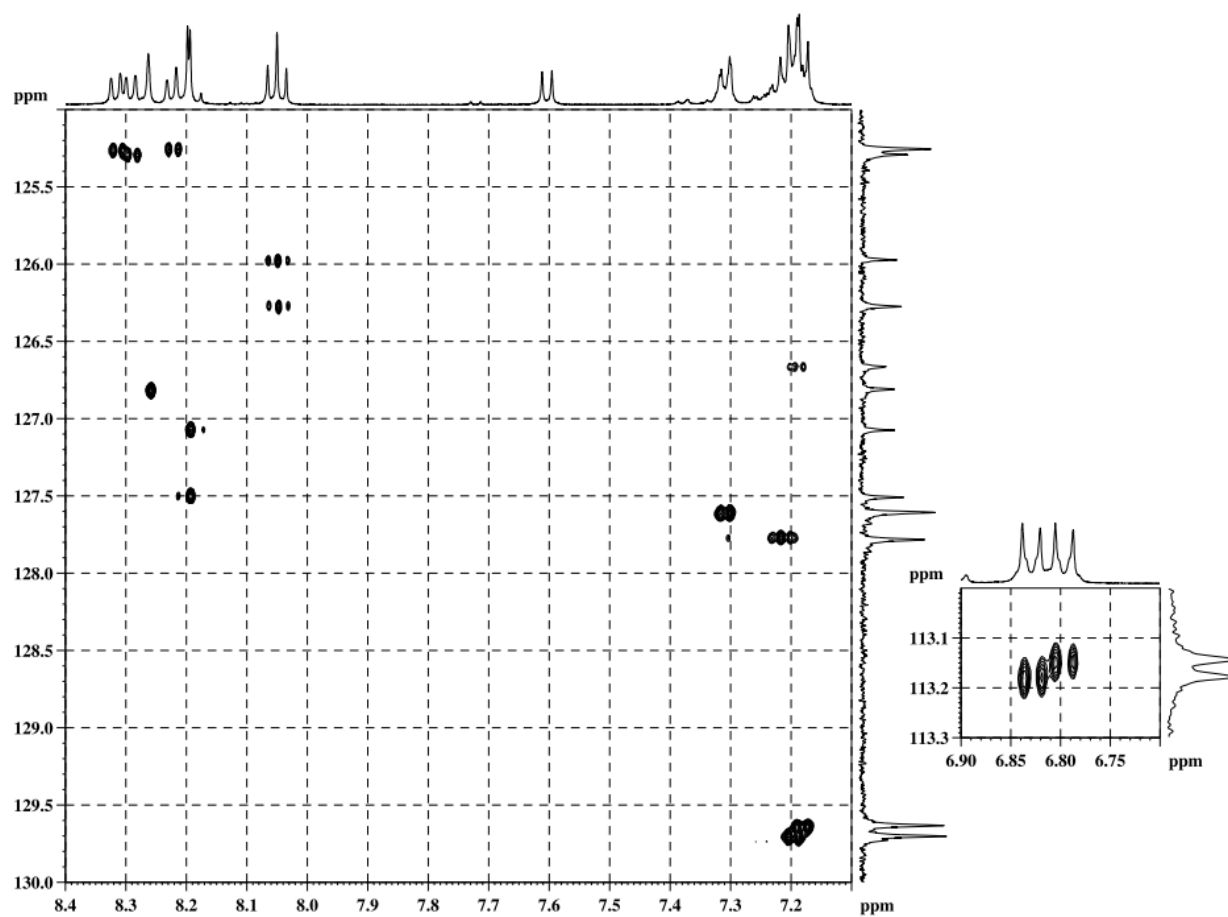


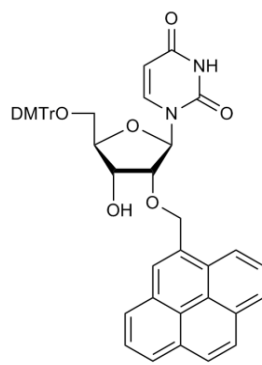




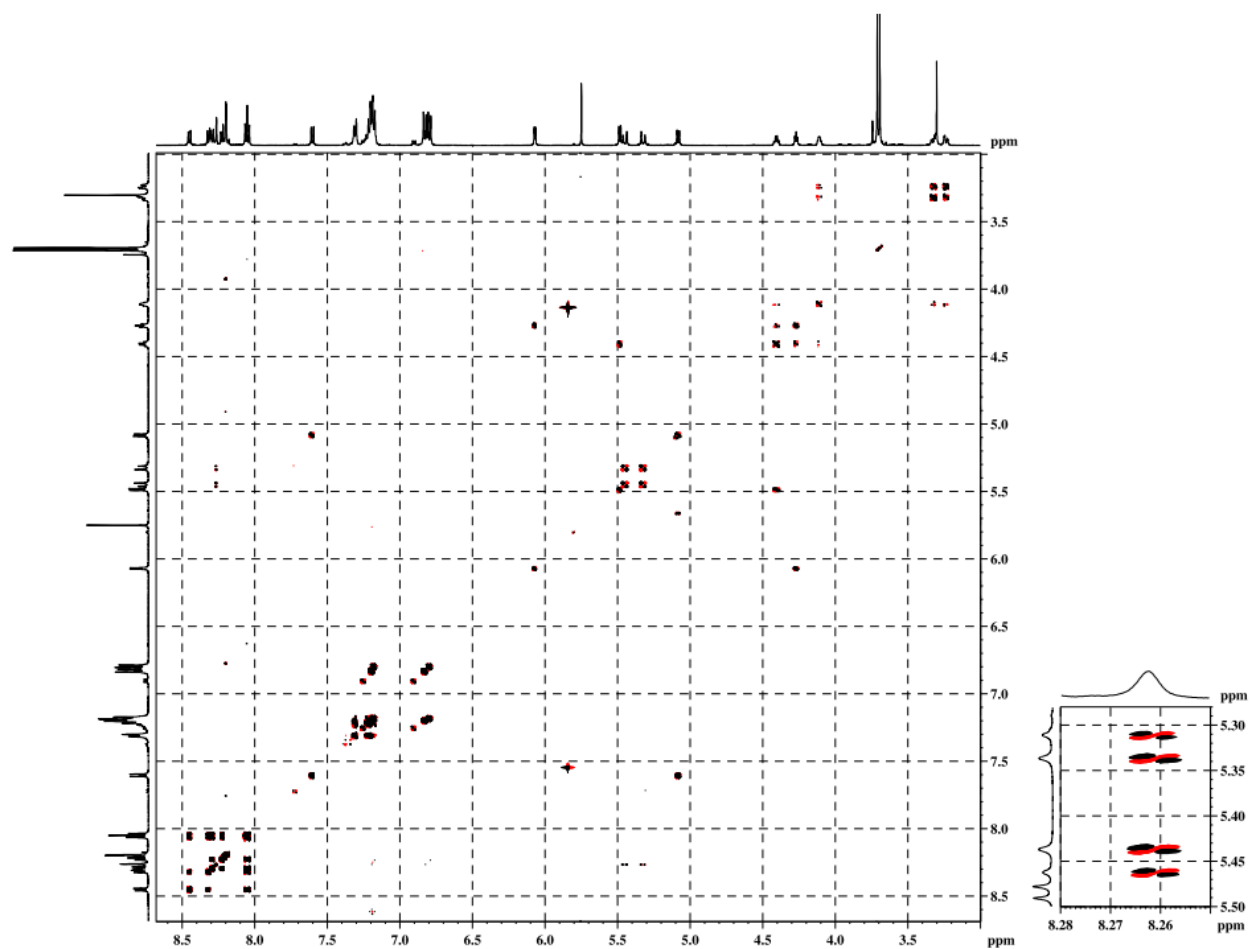


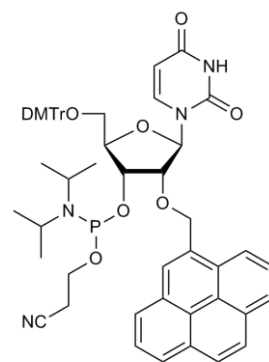
3Z





3Z





4Z

